

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:13:35 ; Search time 1515 Seconds  
(without alignments)  
270.031 Million cell updates/sec

Title: US-09-335-032-71  
Perfect score: 10  
Sequence: 1 cttctctttt 10

Scoring table: OLIGO NUC  
Gapop 60.0 , Gapext 60.0

Searched: 2888711 seqs, 2045481386 residues

Word size : 0

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database : GenEmbl.\*

- 1: gb.ba.\*
- 2: gb.htg.\*
- 3: gb.in.\*
- 4: gb.om.\*
- 5: gb.ov.\*
- 6: gb.pat.\*
- 7: gb.ph.\*
- 8: gb.pl.\*
- 9: gb.pr.\*
- 10: gb.ro.\*
- 11: gb.sts.\*
- 12: gb.sy.\*
- 13: gb.un.\*
- 14: gb.vi.\*
- 15: em.ba.\*
- 16: em.fun.\*
- 17: em.hum.\*
- 18: em.in.\*
- 19: em.mu.\*
- 20: em.om.\*
- 21: em.or.\*
- 22: em.ov.\*
- 23: em.pat.\*
- 24: em.ph.\*
- 25: em.pl.\*
- 26: em.ro.\*
- 27: em.sts.\*
- 28: em.un.\*
- 29: em.vi.\*
- 30: em.htg.hum.\*
- 31: em.htg.inv.\*
- 32: em.htg.other.\*
- 33: em.htg.mus.\*
- 34: em.htg.pln.\*
- 35: em.htg.rod.\*
- 36: em.htg.man.\*
- 37: em.htg.vrt.\*
- 38: em.sy.\*
- 39: em.htgo.hum.\*
- 40: em.htgo.mus.\*
- 41: em.htgo.other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	10	100.0	10	6	BD065135	Character
C 2	10	100.0	12	6	AR029992	Sequence
C 3	10	100.0	12	6	AR030057	Sequence
C 4	10	100.0	17	6	AX500413	Sequence
C 5	10	100.0	17	6	AX500414	Sequence
C 6	10	100.0	17	6	AX500415	Sequence
C 7	10	100.0	17	6	AX500416	Sequence
C 8	10	100.0	17	6	AX500417	Sequence
C 9	10	100.0	17	6	AX500418	Sequence
C 10	10	100.0	17	6	AX500419	Sequence
C 11	10	100.0	17	6	AX500420	Sequence
C 12	10	100.0	17	6	AX673153	Sequence
C 13	10	100.0	19	6	AX659410	Sequence
C 14	10	100.0	20	6	AR315845	Sequence
C 15	10	100.0	20	6	I48976	Sequence 3
C 16	10	100.0	21	6	AR054595	Sequence
C 17	10	100.0	21	6	AR136775	Sequence
C 18	10	100.0	21	6	E35992	Method for
C 19	10	100.0	22	6	AR029876	Sequence
C 20	10	100.0	22	6	AX352316	Sequence
C 21	10	100.0	22	6	AX352317	Sequence
C 22	10	100.0	22	6	AX352318	Sequence
C 23	10	100.0	24	6	AR069183	Sequence
C 24	10	100.0	24	6	AR102693	Sequence
C 25	10	100.0	24	6	AR175558	Sequence
C 26	10	100.0	24	6	AX443310	Sequence
C 27	10	100.0	24	6	BD090389	Sequence
C 28	10	100.0	24	6	BD176467	Method
C 29	10	100.0	24	6	I64401	Sequence 17
C 30	10	100.0	25	6	AX502410	Sequence
C 31	10	100.0	25	6	AX502411	Sequence
C 32	10	100.0	25	6	AX502412	Sequence
C 33	10	100.0	25	6	AX502413	Sequence
C 34	10	100.0	25	6	AX502414	Sequence
C 35	10	100.0	25	6	AX502415	Sequence
C 36	10	100.0	25	6	AX502416	Sequence
C 37	10	100.0	25	6	AX502417	Sequence
C 38	10	100.0	25	6	AX502418	Sequence
C 39	10	100.0	25	6	AX502419	Sequence
C 40	10	100.0	25	6	AX502420	Sequence
C 41	10	100.0	25	6	AX502421	Sequence
C 42	10	100.0	25	6	AX502422	Sequence
C 43	10	100.0	25	6	AX502423	Sequence
C 44	10	100.0	25	6	AX502424	Sequence
C 45	10	100.0	25	6	AX502425	Sequence
C 46	10	100.0	25	6	I48977	Sequence 4
C 47	10	100.0	26	8	ATH525492	Arabidops
C 48	10	100.0	26	8	ATH525503	Arabidops
C 49	10	100.0	27	6	AR066258	Sequence
C 50	10	100.0	27	6	AR191561	Sequence
C 51	10	100.0	27	6	AX115839	Sequence
C 52	10	100.0	28	6	AX040823	Sequence
C 53	10	100.0	29	6	AR261655	Sequence
C 54	10	100.0	30	6	AR261654	Sequence
C 55	10	100.0	30	6	AR261656	Sequence
C 56	10	100.0	30	6	AR261657	Sequence
C 57	10	100.0	33	6	AR120483	Sequence
C 58	10	100.0	33	6	BD063492	Sequence
C 59	10	100.0	33	6	E36416	dna G. 6/20
C 60	10	100.0	38	6	AX218433	Sequence
C 61	10	100.0	38	6	AX220343	Sequence
C 62	10	100.0	38	6	AX273375	Sequence
C 63	10	100.0	40	6	BD182426	Human art
C 64	10	100.0	42	6	I12092	Sequence 33
C 65	10	100.0	43	6	AX484623	Sequence

c 66	10	100.0	45	6	AX061875	Sequence	AX061875	Sequence	10	100.0	111	11	BX295946	Arabidops
c 67	10	100.0	47	8	ATH528602	Arabidops	AU284713	Sequence	10	100.0	112	9	HS1087R	H.sapiens C
c 68	10	100.0	47	6	AR284713	Sequence	AR284713	Sequence	10	100.0	113	8	ATH526836	Arabidops
c 69	10	100.0	47	6	AR288787	Sequence	AR288787	Sequence	10	100.0	113	8	ATH526837	Arabidops
c 70	10	100.0	47	6	AR289547	Sequence	AR289547	Sequence	10	100.0	116	9	HS197G4R	H.sapiens C
c 71	10	100.0	47	6	AR289811	Sequence	AR289811	Sequence	10	100.0	117	9	HS68D7R	H.sapiens C
c 72	10	100.0	47	6	AR290737	Sequence	AR290737	Sequence	10	100.0	117	9	HS68E6F	H.sapiens C
c 73	10	100.0	47	6	AR291549	Sequence	AR291549	Sequence	10	100.0	118	6	AX554775	Sequence
c 74	10	100.0	50	6	AR139658	Sequence	AR139658	Sequence	10	100.0	118	11	HSPE39G05	AL009929 H.sapiens
c 75	10	100.0	50	6	AR158156	Sequence	AR158156	Sequence	10	100.0	120	9	HUMIFARE	M11287 Human alpha
c 76	10	100.0	51	6	AX118141	Sequence	AX118141	Sequence	10	100.0	120	11	HUMSWX752	L24631 Human chrom
c 77	10	100.0	51	6	AX158155	Sequence	AX158155	Sequence	10	100.0	121	6	AX262806	Sequence
c 78	10	100.0	51	6	AX160381	Sequence	AX160381	Sequence	10	100.0	121	6	AX262807	Sequence
c 79	10	100.0	51	6	AX160382	Sequence	AX160382	Sequence	10	100.0	121	6	AX262822	Sequence
c 80	10	100.0	51	6	AX160383	Sequence	AX160383	Sequence	10	100.0	121	6	AX262823	Sequence
c 81	10	100.0	51	6	AX165562	Sequence	AX165562	Sequence	10	100.0	121	6	AX262926	Sequence
c 82	10	100.0	51	8	ATH521150	Arabidops	AJ521150	Arabidops	10	100.0	121	6	AX262930	Sequence
c 83	10	100.0	53	6	AR098682	Sequence	AR098682	Sequence	10	100.0	121	6	AX262931	Sequence
c 84	10	100.0	53	6	AR098683	Sequence	AR098683	Sequence	10	100.0	121	8	AY201631	Arabidops
c 85	10	100.0	53	6	AR204756	Sequence	AR204756	Sequence	10	100.0	121	11	G32506	G32506 A003F19 Hum
c 86	10	100.0	53	6	AR204757	Sequence	AR204757	Sequence	10	100.0	121	11	G32708	G32708 A009Q27 Hum
c 87	10	100.0	54	6	AR134108	Sequence	AR134108	Sequence	10	100.0	122	11	G43213	G43213 WIAF-1715-S
c 88	10	100.0	54	6	AR134285	Sequence	AR134285	Sequence	10	100.0	122	11	G43214	G43214 WIAF-1716-S
c 89	10	100.0	57	8	ATH505724	Arabidops	AJ505724	Arabidops	10	100.0	125	6	BD033545	Sequence
c 90	10	100.0	58	8	ATH527067	Arabidops	AJ527067	Arabidops	10	100.0	128	1	HEA71U	B033452 H.influenza
c 91	10	100.0	61	6	AX270701	Sequence	AX270701	Sequence	10	100.0	128	4	OAU35059	U35059 Ovis aries
c 92	10	100.0	61	6	AX272232	Sequence	AX272232	Sequence	10	100.0	129	11	G32708	G32708 A009Q27 Hum
c 93	10	100.0	65	6	AX482835	Sequence	AX482835	Sequence	10	100.0	130	8	ATH505656	Arabidops
c 94	10	100.0	65	6	AX482852	Sequence	AX482852	Sequence	10	100.0	130	8	ATH528046	Arabidops
c 95	10	100.0	65	6	AX485490	Sequence	AX485490	Sequence	10	100.0	130	9	HS191F10R	AJ528046 Arabidops
c 96	10	100.0	70	6	AR1696	Sequence 1	AR1696	Sequence 1	10	100.0	130	9	HS191F9F	257722 H.sapiens C
c 97	10	100.0	70	6	AR207788	Sequence	AR207788	Sequence	10	100.0	130	11	GL8970	265038 H.sapiens C
c 98	10	100.0	71	6	IG2432	Sequence 5	IG2432	Sequence 5	10	100.0	131	11	HUMSWX53	GL8970 cow STS BMS
c 99	10	100.0	71	9	AH006998S06	Homo sapi	AF026858	Homo sapi	10	100.0	131	11	HUMSWX53	LI4998 Human chrom
c 100	10	100.0	73	6	BD055583	Sequence	BD055583	Sequence	10	100.0	134	9	HSPAI3A8	Z78634 H.sapiens f
c 101	10	100.0	75	6	IG2433	Sequence 6	IG2433	Sequence 6	10	100.0	135	8	AF503161	AF503161 Ceiba pen
c 102	10	100.0	76	6	AX463283	Sequence	AX463283	Sequence	10	100.0	135	10	MUSNR2C06	L35019 Mouse N-met
c 103	10	100.0	76	6	BD038896	Sequence	BD038896	Sequence	10	100.0	136	6	AX135409	AX135409 Sequence
c 104	10	100.0	78	6	IG2446	Sequence 19	IG2446	Sequence 19	10	100.0	136	11	HSP556A10	AL034100 H.sapiens
c 105	10	100.0	80	11	BX000934	Arabidops	BX000934	Arabidops	10	100.0	138	6	BD048886	BD048886 Sequence
c 106	10	100.0	81	14	AB015322	Hepatitis	AB015322	Hepatitis	10	100.0	138	6	BD061135	BD061135 Secreted
c 107	10	100.0	81	14	AF148872	Normal	AF148872	Normal	10	100.0	138	9	HSPAI5A11	Z78713 H.sapiens f
c 108	10	100.0	81	14	AF148873	Normal	AF148873	Normal	10	100.0	138	11	HUMUT799A	L39147 Human STS U
c 109	10	100.0	82	6	IG2447	Sequence 20	IG2447	Sequence 20	10	100.0	140	9	HUMHELAD	D45429 Homo sapien
c 110	10	100.0	83	6	AX386001	Sequence	AX386001	Sequence	10	100.0	140	11	G20401	G20401 human STS A
c 111	10	100.0	83	6	BD037643	Sequence	BD037643	Sequence	10	100.0	141	6	BD055897	BD055897 Sequence
c 112	10	100.0	83	6	BD112411	EST and e	BD112411	EST and e	10	100.0	141	11	AU046839	AU046839 Rattus no
c 113	10	100.0	86	6	AX240905	Sequence	AX240905	Sequence	10	100.0	144	11	AU046839	AU046839 Rattus no
c 114	10	100.0	86	11	DM173H11S	Sequence	DM173H11S	Sequence	10	100.0	145	9	AY190094	AY190094 Homo sapi
c 115	10	100.0	88	11	BX295958	Arabidops	BX295958	Arabidops	10	100.0	146	6	AR139657	AR139657 Sequence
c 116	10	100.0	88	11	G66316	YAC	G66316	YAC	10	100.0	146	11	G02302	G02302 human STS S
c 117	10	100.0	91	3	DME426842	Sequence	DME426842	Sequence	10	100.0	147	3	AF320603	AF320603 Trigon a c
c 118	10	100.0	93	6	AR126785	Sequence	AR126785	Sequence	10	100.0	147	6	AX379108	AX379108 Sequence
c 119	10	100.0	93	6	AR202442	Sequence	AR202442	Sequence	10	100.0	147	6	BD046874	BD046874 Sequence
c 120	10	100.0	93	6	AX522948	Sequence	AX522948	Sequence	10	100.0	147	9	AF515842	AF515842 Homo sapi
c 121	10	100.0	95	6	BD118772	EST and e	BD118772	EST and e	10	100.0	148	6	BD043676	BD043676 Sequence
c 122	10	100.0	98	10	YSCP1PRA	Sequence	YSCP1PRA	Sequence	10	100.0	148	6	BD043896	BD043896 Sequence
c 123	10	100.0	99	6	BD038421	Sequence	BD038421	Sequence	10	100.0	148	6	BD043896	BD043896 Sequence
c 124	10	100.0	99	6	BD038552	Sequence	BD038552	Sequence	10	100.0	148	9	AY190090	AY190090 Homo sapi
c 125	10	100.0	99	6	BD038552	Sequence	BD038552	Sequence	10	100.0	148	9	HUMFCG8A02	M90722 Human FC-ga
c 126	10	100.0	102	11	G20394	human STS A	G20394	human STS A	10	100.0	148	9	HUMFCG8A02	M90722 Human FC-ga
c 127	10	100.0	103	8	OSA532497	Sequence	OSA532497	Sequence	10	100.0	149	6	AX259898	AX259898 Sequence
c 128	10	100.0	103	9	HSSTEA4TF1	Sequence	HSSTEA4TF1	Sequence	10	100.0	149	6	AX259898	AX259898 Sequence
c 129	10	100.0	103	11	G19611	human STS A	G19611	human STS A	10	100.0	150	11	G59542	Z61827 H.sapiens C
c 130	10	100.0	104	6	AX645544	Sequence	AX645544	Sequence	10	100.0	150	11	G59725	G59725 SHGC-130478
c 131	10	100.0	104	6	AX676705	Sequence	AX676705	Sequence	10	100.0	151	9	AY190102	AY190102 Homo sapi
c 132	10	100.0	104	6	BD049479	Sequence	BD049479	Sequence	10	100.0	151	6	BD028580	BD028580 Sequence
c 133	10	100.0	104	8	AY220694	Sequence	AY220694	Sequence	10	100.0	152	6	BD028580	BD028580 Sequence
c 134	10	100.0	105	6	AX127420	Sequence	AX127420	Sequence	10	100.0	152	9	AY190114	AY190114 Homo sapi
c 135	10	100.0	105	11	BX248221	Arabidops	BX248221	Arabidops	10	100.0	152	9	HUMALURPTF	M37552 Human AFP g
c 136	10	100.0	105	11	G71016	Sequence	G71016	Sequence	10	100.0	153	5	AY269268	AY269268 Gila eleg
c 137	10	100.0	107	9	HUMHELAC	Sequence	HUMHELAC	Sequence	10	100.0	153	5	AX269268	AX269268 Gila eleg
c 138	10	100.0	110	11	BX295578	Arabidops	BX295578	Arabidops	10	100.0	153	9	HSXPA44B	X92876 H.sapiens m



C 358	10	100.0	217	6	BD030815	BD030815 Sequence	431	10	100.0	236	8	AB043752	AB043752 Fagus cre
C 359	10	100.0	217	6	BD055876	BD055876 Sequence	432	10	100.0	236	8	AB043753	AB043753 Fagus cre
C 360	10	100.0	217	9	HSOBR128	U62518 Human ORF g	433	10	100.0	236	8	AB043757	AB043757 Fagus cre
C 361	10	100.0	217	11	G04642	G04642 human STS W	434	10	100.0	236	8	AB043758	AB043758 Fagus cre
C 362	10	100.0	218	6	BD056063	BD056063 Sequence	435	10	100.0	236	8	AB043759	AB043759 Fagus cre
C 363	10	100.0	218	9	HSTNFR2507	U52162 Human tumor	436	10	100.0	236	8	AB043760	AB043760 Fagus jap
C 364	10	100.0	218	10	MMU409500	AJ409500 Mus muscu	437	10	100.0	236	8	AB043761	AB043761 Fagus jap
C 365	10	100.0	219	1	STRMLIKEB	L05018 Streptococc	438	10	100.0	236	8	AB043762	AB043762 Fagus jap
C 366	10	100.0	219	3	AF323685	Z55009 H. sapiens C	439	10	100.0	236	8	AB066498	AB066498 Fagus gra
C 367	10	100.0	219	9	HS184A2F	Z55009 H. sapiens C	440	10	100.0	236	8	AB070965	AB070965 Fagus syl
C 368	10	100.0	219	11	ASP217F08	AL0009571 H. sapiens	441	10	100.0	236	8	AY022125	AY022125 Oryza sat
C 369	10	100.0	220	8	HSP00757	K02267 Human enkep	442	10	100.0	236	8	AY022309	AY022309 Oryza sat
C 370	10	100.0	221	9	HUMENKB1	G35200 cre12e Cryp	443	10	100.0	237	1	AF093626	AF093626 Bacillus
C 371	10	100.0	221	11	G35200	AU05857 Homo sapi	444	10	100.0	237	5	GGZ94841	Z94841 G.gallus mi
C 372	10	100.0	222	9	HS405857	L34731 Homo sapien	445	10	100.0	237	9	HS55G9R	Z65737 H.sapiens C
C 373	10	100.0	222	9	HUMTCRBA1	L34736 Homo sapien	446	10	100.0	237	9	HSU39128	U93128 Homo sapien
C 374	10	100.0	222	9	HUMTCRBA1	U96350 Hepatitis G	447	10	100.0	237	9	HUMDINUCRP	M94958 Human dinuc
C 375	10	100.0	222	14	U96350	BD032653 Sequence	448	10	100.0	238	6	AX094172	AX094172 Sequence
C 376	10	100.0	223	6	BD032653	AY032840 Passiflor	449	10	100.0	238	6	AX094172	AX094172 Sequence
C 377	10	100.0	223	6	AY032840	G26013 human STS E	450	10	100.0	238	11	G63246	G63246 SHGC-140885
C 378	10	100.0	223	11	G26013	U96357 Hepatitis G	451	10	100.0	239	5	GGZ94829	Z94829 G.gallus mi
C 379	10	100.0	223	14	U96357	AY017798 Oryza sat	452	10	100.0	239	6	BD028893	BD028893 Sequence
C 380	10	100.0	224	8	AY017798	AY020653 Oryza sat	453	10	100.0	239	6	BD048305	BD048305 Sequence
C 381	10	100.0	224	8	AY020653	AY023853 Oryza sat	454	10	100.0	239	6	BD059663	BD059663 Secreted
C 382	10	100.0	224	8	AY023853	BD033343 Sequence	455	10	100.0	239	8	AB084875	AB084875 Schizosac
C 383	10	100.0	225	6	BD033343	AF374108 Fusarium	456	10	100.0	239	11	G04386	G04386 human STS W
C 384	10	100.0	225	8	AF374108	AY077694 Cantharel	457	10	100.0	240	6	BD071286	BD071286 Secreted
C 385	10	100.0	225	9	HS30A2R	Z55278 H. sapiens C	458	10	100.0	240	8	AY021176	AY021176 Oryza sat
C 386	10	100.0	225	8	AY077694	BD071601 Secreted	459	10	100.0	240	9	AF041219	AF041219 Homo sapi
C 387	10	100.0	225	9	HS4031790	AJ403790 Homo sapi	460	10	100.0	240	9	F267517S12	F267517S12 Homo sapi
C 388	10	100.0	226	6	BD071601	AY017828 Oryza sat	461	10	100.0	240	9	HS405839	AJ405839 Homo sapi
C 389	10	100.0	226	8	AY017828	AY017828 Oryza sat	462	10	100.0	240	9	S63249	S63249 T-cell rece
C 390	10	100.0	226	8	AY019941	AY026388 Rattus no	463	10	100.0	240	11	AU025673	AU025673 Rattus no
C 391	10	100.0	226	8	AY019941	AY026388 Rattus no	464	10	100.0	241	5	GGZ95323	Z95323 G.gallus mi
C 392	10	100.0	226	11	AY026388	AU048388 Rattus no	465	10	100.0	241	6	AR240300	AR240300 Sequence
C 393	10	100.0	226	11	AU048388	G02977 human STS W	466	10	100.0	241	6	AR280852	AR280852 Sequence
C 394	10	100.0	226	11	G02977	G05222 human STS W	467	10	100.0	241	6	AR283348	AR283348 Sequence
C 395	10	100.0	226	11	G05222	AJ296842 Homo sapi	468	10	100.0	241	6	AX303040	AX303040 Sequence
C 396	10	100.0	226	11	HS296842	AJ296842 Homo sapi	469	10	100.0	242	8	AY017747	AY017747 Oryza sat
C 397	10	100.0	227	8	AB043754	AB043754 Fagus cre	470	10	100.0	242	9	HS46F12R	Z58627 H. sapiens C
C 398	10	100.0	227	8	AY023637	AY023637 Oryza sat	471	10	100.0	243	6	AX048534	AX048534 Sequence
C 399	10	100.0	227	8	AY023637	U01283 Rattus norv	472	10	100.0	243	11	G36317	G36317 STS hl4250
C 400	10	100.0	227	11	RNM3DDE03	G05891 human STS W	473	10	100.0	244	9	HS13B3F	Z56737 H. sapiens C
C 401	10	100.0	227	11	G05891	AR269536 Sequence	474	10	100.0	245	8	AF374129	AF374129 Fusarium
C 402	10	100.0	228	6	AR269536	AX437220 Sequence	475	10	100.0	245	9	HS403843	AJ403843 Homo sapi
C 403	10	100.0	228	6	AX437220	AF076728 Dendrochi	476	10	100.0	245	11	AL823657	AL823657 Arabidops
C 404	10	100.0	228	8	AF076728	AY018284 Oryza sat	477	10	100.0	246	6	AX185506	Z52141 H. sapiens C
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C 409	10	100.0	228	11	HS405793	G67231 csnpkU86-pc	482	10	100.0	246	11	G49085	G49085 SHGC-83039
C 410	10	100.0	228	11	G67231	X51790 Human mRNA	483	10	100.0	247	6	AR270174	AR270174 Sequence
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C 412	10	100.0	229	11	AU027640	AY022834 Oryza sat	485	10	100.0	247	6	BD043817	BD043817 Kluyverom
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C 417	10	100.0	231	8	HFI3111824	X51788 Human mRNA	490	10	100.0	249	8	AF374105	AF374105 Fusarium
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C 419	10	100.0	232	6	BD117604	AF374120 Fusarium	492	10	100.0	249	8	AF374109	AF374109 Fusarium
C 420	10	100.0	232	8	AF374120	BD139321 Extended	493	10	100.0	249	8	AF374110	AF374110 Fusarium
C 421	10	100.0	232	6	BD139321	AJ389986 Homo sapi	494	10	100.0	249	8	AF374111	AF374111 Fusarium
C 422	10	100.0	233	9	HS4389986	X51794 Human mRNA	495	10	100.0	249	8	AF374112	AF374112 Fusarium
C 423	10	100.0	233	9	HSCTL67	G13001 SNSS2840 Er	496	10	100.0	249	8	AF374113	AF374113 Fusarium
C 424	10	100.0	233	11	G13001	X17075 Caenorhabdi	497	10	100.0	249	8	AF374114	AF374114 Fusarium
C 425	10	100.0	234	3	CE3	AJ405685 Homo sapi	498	10	100.0	249	8	AF374115	AF374115 Fusarium
C 426	10	100.0	234	9	HS4405685	Z94830 G.gallus mi	499	10	100.0	249	8	AF374116	AF374116 Fusarium
C 427	10	100.0	235	5	GGZ94830	AF308440 Mus muscu	500	10	100.0	249	8	AF374117	AF374117 Fusarium
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ALIGNMENTS



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RESULT 1
BD065135          10 bp      DNA      linear      PAT 27-AUG-2002
LOCUS
DEFINITION      Characterization of the yeast transcriptome.
ACCESSION      BD065135
VERSION        BD065135.1 GI:22610738
KEYWORDS
SOURCE        Saccharomyces cerevisiae (baker's yeast)
ORGANISM
REFERENCE
AUTHORS        Wang, C.-G. and Hepburn, A.G.
TITLE          Genetic sequence assay using DNA triple strand formation
JOURNAL        Patent: US 5861244-A 71 10-JUL-2001;
              THE JOHNS HOPKINS UNIVERSITY SCHOOL OF MEDICINE
COMMENT
PN JP 2001509017-A/71
PD 10-JUL-2001
PF 22-JAN-1998 JP 1998532117
PR 23-JAN-1997 US 60/035917
PI VICTOR E VELCULESCU, BERT VOGELSTEIN, KENNETH W KINZLER PC
CI 2N15/10, CI 2N15/31, C07K14/395, C12Q1/68, C12Q1/02 CC
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FT Location/Qualifiers
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DEFINITION      Sequence 181 from patent US 5861244.
ACCESSION      AR029992
VERSION        AR029992.1 GI:5943206
KEYWORDS
SOURCE        Unknown.
ORGANISM
REFERENCE
AUTHORS        Wang, C.-G. and Hepburn, A.G.
TITLE          Genetic sequence assay using DNA triple strand formation
JOURNAL        Patent: US 5861244-A 191 19-JAN-1999;
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Db 11 CTCTCTCTTTT 2

RESULT 3
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LOCUS
DEFINITION      Characterization of the yeast transcriptome.
ACCESSION      BD065135
VERSION        BD065135.1 GI:22610738
KEYWORDS
SOURCE        Saccharomyces cerevisiae (baker's yeast)
ORGANISM
REFERENCE
AUTHORS        Wang, C.-G. and Hepburn, A.G.
TITLE          Genetic sequence assay using DNA triple strand formation
JOURNAL        Patent: US 5861244-A 71 10-JUL-2001;
              THE JOHNS HOPKINS UNIVERSITY SCHOOL OF MEDICINE
COMMENT
PN JP 2001509017-A/71
PD 10-JUL-2001
PF 22-JAN-1998 JP 1998532117
PR 23-JAN-1997 US 60/035917
PI VICTOR E VELCULESCU, BERT VOGELSTEIN, KENNETH W KINZLER PC
CI 2N15/10, CI 2N15/31, C07K14/395, C12Q1/68, C12Q1/02 CC
CH Characterization of the yeast transcriptome
FH Key
FT source
FT Location/Qualifiers
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source
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RESULT 4
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LOCUS
DEFINITION      Sequence 1720 from Patent EP1229046.
ACCESSION      AX500413
VERSION        AX500413.1 GI:23382706
KEYWORDS
SOURCE        Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS        Zhan, J.
TITLE          Human testis expressed patched like protein
JOURNAL        Patent: EP 1229046-A 1720 07-AUG-2002;
              Aeomica, Inc. (US)
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DEFINITION      Sequence 1721 from Patent EP1229046.
ACCESSION      AX500414
VERSION        AX500414.1 GI:23382707
KEYWORDS
SOURCE        Homo sapiens (human)
ORGANISM

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RESULT 3
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DEFINITION      Sequence 246 from patent US 5861244.
ACCESSION      AR030057
VERSION        AR030057.1 GI:5943271
KEYWORDS
SOURCE        Unknown.
ORGANISM
REFERENCE
AUTHORS        Wang, C.-G. and Hepburn, A.G.
TITLE          Genetic sequence assay using DNA triple strand formation
JOURNAL        Patent: US 5861244-A 246 19-JAN-1999;
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LOCUS
DEFINITION      Sequence 1720 from Patent EP1229046.
ACCESSION      AX500413
VERSION        AX500413.1 GI:23382706
KEYWORDS
SOURCE        Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS        Zhan, J.
TITLE          Human testis expressed patched like protein
JOURNAL        Patent: EP 1229046-A 1720 07-AUG-2002;
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Db 17 CTCTCTCTTTT 8

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DEFINITION      Sequence 1721 from Patent EP1229046.
ACCESSION      AX500414
VERSION        AX500414.1 GI:23382707
KEYWORDS
SOURCE        Homo sapiens (human)
ORGANISM

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
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Human testis expressed patched like protein  
Patent: EP 1229046-A 1721 07-AUG-2002;  
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Db 16 CTTCTCTTTT 7  
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LOCUS AX500415 17 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 1722 from Patent EP1229046.  
ACCESSION AX500415  
VERSION AX500415.1 GI:23382708  
KEYWORDS Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
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Human testis expressed patched like protein  
Patent: EP 1229046-A 1722 07-AUG-2002;  
Aeomica, Inc. (US)  
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LOCUS AX500416 17 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 1723 from Patent EP1229046.  
ACCESSION AX500416  
VERSION AX500416.1 GI:23382709  
KEYWORDS Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
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Human testis expressed patched like protein  
Patent: EP 1229046-A 1723 07-AUG-2002;  
Aeomica, Inc. (US)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
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Human testis expressed patched like protein  
Patent: EP 1229046-A 1721 07-AUG-2002;  
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DEFINITION Sequence 1724 from Patent EP1229046.  
ACCESSION AX500417  
VERSION AX500417.1 GI:23382710  
KEYWORDS Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
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Human testis expressed patched like protein  
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LOCUS AX500418 17 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 1725 from Patent EP1229046.  
ACCESSION AX500418  
VERSION AX500418.1 GI:23382711  
KEYWORDS Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
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Human testis expressed patched like protein  
Patent: EP 1229046-A 1725 07-AUG-2002;  
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LOCUS      AX500419      17 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 1726 from Patent EP1229046.
ACCESSION  AX500419
VERSION     AX500419.1 GI:23382712
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1
AUTHORS     Zhan,J.
TITLE       Human testis expressed patched like protein
JOURNAL     Patent: EP 1229046-A 1726 07-AUG-2002;
            Aescima, Inc. (US)
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Db 11 CTTCTCTTTT 2

RESULT 11
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LOCUS      AX500420      17 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 1727 from Patent EP1229046.
ACCESSION  AX500420
VERSION     AX500420.1 GI:23382713
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1
AUTHORS     Zhan,J.
TITLE       Human testis expressed patched like protein
JOURNAL     Patent: EP 1229046-A 1727 07-AUG-2002;
            Aescima, Inc. (US)
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QY 1 CTTCTCTTTT 10
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RESULT 14

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Db 10 CTTCTCTTTT 1

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LOCUS      AX673153      17 bp      DNA      linear      PAT 27-MAR-2003
DEFINITION Sequence 1598 from Patent WO03004526.
ACCESSION  AX673153
VERSION     AX673153.1 GI:29331501
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1
AUTHORS     Telerman,A., Anson,R. and Tuijnder,M.
TITLE       Sequences involved in phenomena of tumour suppression, tumour
            reversion, apoptosis and/or resistance to viruses and their use as
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JOURNAL     Patent: WO 03004526-A 1598 16-JAN-2003;
            Molecular Engines Laboratories (FR)
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LOCUS      AX659410      19 bp      DNA      linear      PAT 22-MAR-2003
DEFINITION Sequence 12 from Patent WO02102824.
ACCESSION  AX659410
VERSION     AX659410.1 GI:29161640
KEYWORDS
SOURCE      synthetic construct
            synthetic construct
            artificial sequences.
REFERENCE   1
AUTHORS     Beifehr,C. and Snaidr,J.
TITLE       Method for specific fast detection of relevant bacteria in drinking
            water
JOURNAL     Patent: WO 02102824-A 12 27-DEC-2002;
            Vermicon AG (DE)
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QY 1 CTTCTCTTTT 10
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RESULT 14

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AR315845/c
LOCUS AR315845 20 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 6382 from patent US 6559294.
ACCESSION AR315845
VERSION AR315845.1 GI:31709271
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Griffais, R., Hoigeth, S.K., Zagursky, R.J., Metcalf, B.J., Peek, J.A.,
Sankaran, B., and Fletcher, L.D.
TITLE Chlamydia pneumoniae polynucleotides and uses thereof
JOURNAL Patent: US 6559294-A 6382 06-MAY-2003;
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Qy 1 CTTCTCTTTT 10
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RESULT 15
LOCUS I48976 20 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 3 from patent US 5627054.
ACCESSION I48976
VERSION I48976.1 GI:2467439
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Gillespie, D. deceased.
TITLE Competitor primer asymmetric polymerase chain reaction
JOURNAL Patent: US 5627054-A 3 06-MAY-1997;
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Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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RESULT 16
LOCUS AR054595/c 21 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 16 from patent US 5837447.
ACCESSION AR054595
VERSION AR054595.1 GI:5980172
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Gorski, J.
TITLE Monitoring an immune response by analysis of amplified
immunoglobulin or T-cell-receptor nucleic acid
JOURNAL Patent: US 5837447-A 16 17-NOV-1998;

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RESULT 17
LOCUS ARI36775/c 21 bp DNA linear PAT 16-JUN-2001
DEFINITION Sequence 8 from patent US 6162435.
ACCESSION ARI36775
VERSION ARI36775.1 GI:14478025
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Minion, F. Chris. and Hsu, T.
TITLE Recombinant mycoplasma hyopneumoniae vaccine
JOURNAL Patent: US 6162435-A 8 19-DEC-2000;
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BASE COUNT 12 a 0 c 7 g 2 t
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RESULT 18
LOCUS E35992/c 21 bp DNA linear PAT 31-JAN-2002
DEFINITION Method for detecting Kawasaki disease factor.
ACCESSION E35992
VERSION E35992.1 GI:18624703
KEYWORDS JP 2000157297-A/83.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 21)
AUTHORS Yoshioka, T. and Suzuki, R.
TITLE Method for detecting Kawasaki disease factor
JOURNAL Patent: JP 2000157297-A 83 13-JUN-2000;
COMMENT SHIONOGI & CO LTD
OS Artificial Sequence
PN JP 2000157297-A/83
PD 13-JUN-2000
PF 01-DEC-1998 JP 1998341661
PR TAKESHI YOSHIOKA, RYUJI SUZUKI
PI C12Q1/68, C12N15/09, G01N33/48, C12N15/00
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/organism="Artificial Sequence".
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source Location/Qualifiers
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/organism="synthetic construct"

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BASE COUNT      8 a      3 c      7 g      3 t
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Query Match      100.0%; Score 10; DB 6; Length 21;
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Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
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Db 18 CTTCTCTTTT 9

RESULT 19
LOCUS      AR029876      22 bp      DNA      linear      PAT 29-SEP-1999
DEFINITION      Sequence 65 from patent US 5861244.
ACCESSION      AR029876
VERSION      AR029876.1 GI:5943090
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      1 (bases 1 to 22)
AUTHORS      Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL      Patent: US 5861244-A 65 19-JAN-1999;
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Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
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Db 6 CTTCTCTTTT 15

RESULT 20
LOCUS      AX352316      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 612 from Patent WO0193902.
ACCESSION      AX352316
VERSION      AX352316.1 GI:18617599
KEYWORDS
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      1
AUTHORS      Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 612 13-DEC-2001;
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BASE COUNT      1 a      4 c      3 g      14 t
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Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
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Db 1 CTTCTCTTTT 10

RESULT 21
LOCUS      AX352317      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 613 from Patent WO0193902.
ACCESSION      AX352317
VERSION      AX352317.1 GI:18617600
KEYWORDS
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      1
AUTHORS      Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 613 13-DEC-2001;
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BASE COUNT      1 a      4 c      4 g      13 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
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Db 7 CTTCTCTTTT 16

RESULT 22
LOCUS      AX352318      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 614 from Patent WO0193902.
ACCESSION      AX352318
VERSION      AX352318.1 GI:18617601
KEYWORDS
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      1
AUTHORS      Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 614 13-DEC-2001;
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            /mol_type="genomic DNA"
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            /note="Synthetic HDR"
BASE COUNT      1 a      4 c      2 g      15 t
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Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 7 CTTCTCTTTT 16

RESULT 23
LOCUS      AR069183/c      24 bp      DNA      linear      PAT 18-FEB-2000
DEFINITION      Sequence 17 from patent US 5891623.
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ACCESSION      AR069183
VERSION        AR069183.1  GI:7220071
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Primi,D.
TITLE        Diagnosis and treatment of AIDS onset
JOURNAL      Patent: US 5891623-A 17 06-APR-1999;
FEATURES     Location/Qualifiers
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              /organism="unknown"
BASE COUNT   10 a 3 c 7 g 4 t
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Query Match   100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 24
ACCESSION      AR102693/c
LOCUS          AR102693
DEFINITION     Sequence 16 from patent US 6087096.
ACCESSION      AR102693
VERSION        AR102693.1  GI:12814281
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Dau,P.C. and Liu,D.
TITLE        Method of intrafamily fragment analysis of the T cell receptor
JOURNAL      Patent: US 6087096-A 16 11-JUL-2000;
FEATURES     Location/Qualifiers
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              /organism="unknown"
BASE COUNT   10 a 3 c 7 g 4 t
ORIGIN
Query Match   100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 25
ACCESSION      AR175558/c
LOCUS          AR175558
DEFINITION     Sequence 3 from patent US 6309837.
ACCESSION      AR175558
VERSION        AR175558.1  GI:117916857
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Dean,R.A. and Wang,Y.-H.
TITLE        PCR-based method for identifying a fusarium wilt-resistant genotype
JOURNAL      Patent: US 6309837-A 3 30-OCT-2001;
FEATURES     Location/Qualifiers
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              /organism="unknown"
source

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BASE COUNT   14 a 1 c 8 g 1 t
ORIGIN
Query Match   100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 23 CTTCTCTTTT 14

RESULT 26
ACCESSION      AX443310/c
LOCUS          AX443310
DEFINITION     Sequence 46 from Patent WO0216940.
ACCESSION      AX443310
VERSION        AX443310.1  GI:21690705
KEYWORDS
SOURCE        synthetic construct
ORGANISM      synthetic construct
              artificial sequences.
REFERENCE     1
AUTHORS      Sulavik,M., Ling,L.L., Opperman,T., Moir,D.T. and Bunker,C.
TITLE        Genomics-assisted rapid identification of targets
JOURNAL      Patent: WO 0216940-A 46 28-FEB-2002;
              Genome Therapeutics Corporation (US)
FEATURES     Location/Qualifiers
              1..24
              /organism="synthetic construct"
              /mol_type="genomic DNA"
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              /note="primer"
BASE COUNT   9 a 4 c 6 g 5 t
ORIGIN
Query Match   100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 16 CTTCTCTTTT 7

RESULT 27
ACCESSION      BD090389/c
LOCUS          BD090389
DEFINITION     A method of arraying genome clone.
ACCESSION      BD090389
VERSION        BD090389.1  GI:22635999
KEYWORDS
SOURCE        synthetic construct
ORGANISM      synthetic construct
              artificial sequences.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Soeda,E.
TITLE        A method of arraying genome clone
JOURNAL      Patent: JP 2001321190-A 2633 20-NOV-2001;
              THE INSTITUTE OF PHYSICAL AND CHEMICAL RESEARCH, YUGENKAISHA
              GENOTECHS
COMMENT       OS Artificial Sequence
              PN JP 2001321190-A/2633
              PD 20-NOV-2001
              PF 12-MAR-2001 JP 2001068285
              PI EIICHI SOEDA
              PC C12N15/09,C12N15/09,C12M1/00,C12Q1/68,G01N33/53,G01N33/566, PC
              C12N15/00
              CC C12N15/00
              Description of Artificial Sequence:Synthetic DNA FH Key
              Location/Qualifiers
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              /organism='Artificial Sequence'.
              FT source
              FT

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FEATURES
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BASE COUNT    12 a    2 c    6 g    4 t
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Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 10
    |||||

RESULT 28
BD176467/c
LOCUS      BD176467       24 bp    DNA    linear    PAT 18-MAR-2003
DEFINITION A method of arraying genome clone.
ACCESSION  BD176467
VERSION     BD176467.1 GI:29122175
KEYWORDS   WO 02072815-A/267.
SOURCE     synthetic construct
ORGANISM   artificial sequences.
REFERENCE  1 (bases 1 to 24)
AUTHORS    Soeda,E.
TITLE      A method of arraying genome clone
JOURNAL    Patent: WO 02072815-A 267 19-SEP-2002;
            EIIICHI SOEDA,TAKESHI KUKITA
COMMENT    OS Artificial Sequence
            PN WO 02072815-A/267
            PD 19-SEP-2002 WO 2001JP004139
            PF 17-MAY-2001 WO 2001JP004139
            PR 12-MAR-2001 JP 01P 68285
            PT EIIICHI SOEDA
            PC C12N15/09,C12Q1/68
            CC Description of Artificial Sequence: Synthetic DNA FH Key
            FT source
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                 /organism="synthetic construct"
                 /mol_type="genomic DNA"
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BASE COUNT    12 a    2 c    6 g    4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 10
    |||||

RESULT 29
I64401/c
LOCUS      I64401       24 bp    DNA    linear    PAT 07-OCT-1997
DEFINITION Sequence 17 from patent US 5665355.
ACCESSION  I64401
VERSION     I64401.1 GI:2481295
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unknown.
REFERENCE  1 (bases 1 to 24)
AUTHORS    Primi,D.
TITLE      Diagnosis and treatment of AIDS onset

FEATURES
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      /mol_type="genomic DNA"
      /db_xref="taxon:32630"
BASE COUNT    12 a    2 c    6 g    4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 10
    |||||

RESULT 30
AX502410/c
LOCUS      AX502410       25 bp    DNA    linear    PAT 27-SEP-2002
DEFINITION Sequence 3717 from Patent EP1229046.
ACCESSION  AX502410
VERSION     AX502410.1 GI:23384703
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
REFERENCE  1
AUTHORS    Zhan,J.
TITLE      Human testis expressed patched like protein
JOURNAL    Patent: EP 1229046-A 3717 07-AUG-2002;
            Aeomica, Inc. (US)
COMMENT    /organism="Homo sapiens"
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BASE COUNT    9 a    2 c    9 g    5 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
    |||||
Db 25 CTTCTCTTTT 16
    |||||

RESULT 31
AX502411/c
LOCUS      AX502411       25 bp    DNA    linear    PAT 27-SEP-2002
DEFINITION Sequence 3718 from Patent EP1229046.
ACCESSION  AX502411
VERSION     AX502411.1 GI:23384704
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
REFERENCE  1
AUTHORS    Zhan,J.
TITLE      Human testis expressed patched like protein
JOURNAL    Patent: EP 1229046-A 3718 07-AUG-2002;
            Aeomica, Inc. (US)
COMMENT    /organism="Homo sapiens"
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BASE COUNT    8 a    2 c    9 g    6 t
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JOURNAL Patent: US 5665355-A 17 09-SEP-1997;
  source Location/Qualifiers
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BASE COUNT    10 a    3 c    7 g    4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
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Db 18 CTTCTCTTTT 9
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Query Match 100.0%; Score 10; DB 6; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 24 CTTCTCTTTT 15

RESULT 32  
AX502412/c  
LOCUS AX502412 25 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 3719 from Patent EP1229046.  
ACCESSION AX502412  
VERSION AX502412.1 GI:23384705  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1  
AUTHORS Zhan, J.  
TITLE Human testis expressed patched like protein  
JOURNAL Patent: EP 1229046-A 3719 07-AUG-2002;  
Aeomica, Inc. (US)

FEATURES  
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Location/Qualifiers  
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/mol\_type="genomic DNA"  
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BASE COUNT  
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Query Match 100.0%; Score 10; DB 6; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 23 CTTCTCTTTT 14

RESULT 33  
AX502413/c  
LOCUS AX502413 25 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 3720 from Patent EP1229046.  
ACCESSION AX502413  
VERSION AX502413.1 GI:23384706  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1  
AUTHORS Zhan, J.  
TITLE Human testis expressed patched like protein  
JOURNAL Patent: EP 1229046-A 3720 07-AUG-2002;  
Aeomica, Inc. (US)

FEATURES  
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BASE COUNT  
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Query Match 100.0%; Score 10; DB 6; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 22 CTTCTCTTTT 13

RESULT 34  
AX502414/c  
LOCUS AX502414 25 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 3721 from Patent EP1229046.  
ACCESSION AX502414  
VERSION AX502414.1 GI:23384707  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1  
AUTHORS Zhan, J.  
TITLE Human testis expressed patched like protein  
JOURNAL Patent: EP 1229046-A 3721 07-AUG-2002;  
Aeomica, Inc. (US)

FEATURES  
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Location/Qualifiers  
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/mol\_type="genomic DNA"  
/db\_xref="taxon:9606"  
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BASE COUNT  
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Query Match 100.0%; Score 10; DB 6; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 21 CTTCTCTTTT 12

RESULT 35  
AX502415/c  
LOCUS AX502415 25 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 3722 from Patent EP1229046.  
ACCESSION AX502415  
VERSION AX502415.1 GI:23384708  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1  
AUTHORS Zhan, J.  
TITLE Human testis expressed patched like protein  
JOURNAL Patent: EP 1229046-A 3722 07-AUG-2002;  
Aeomica, Inc. (US)

FEATURES  
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BASE COUNT  
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Query Match 100.0%; Score 10; DB 6; Length 25;  
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Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 20 CTTCTCTTTT 11

RESULT 36  
AX502416/c  
LOCUS AX502416 25 bp DNA linear PAT 27-SEP-2002  
DEFINITION Sequence 3723 from Patent EP1229046.  
ACCESSION AX502416



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VERSION AX502416.1 GI:23384709
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3725 07-AUG-2002;
Aeomica, Inc. (US)
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/db_xref="taxon:9606"
BASE COUNT 9 a 1 c 8 g 7 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 19 CTTCTCTTTT 10

RESULT 37
AX502417/c
LOCUS AX502417 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3724 from Patent EP1229046.
ACCESSION AX502417
VERSION AX502417.1 GI:23384710
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3724 07-AUG-2002;
Aeomica, Inc. (US)
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 38
AX502418/c
LOCUS AX502418 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3725 from Patent EP1229046.
ACCESSION AX502418
VERSION AX502418.1 GI:23384711
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3725 07-AUG-2002;
Aeomica, Inc. (US)
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/db_xref="taxon:9606"
BASE COUNT 9 a 2 c 9 g 5 t
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 39
AX502419/c
LOCUS AX502419 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3726 from Patent EP1229046.
ACCESSION AX502419
VERSION AX502419.1 GI:23384712
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3726 07-AUG-2002;
Aeomica, Inc. (US)
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BASE COUNT 9 a 3 c 8 g 5 t
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 40
AX502420/c
LOCUS AX502420 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3727 from Patent EP1229046.
ACCESSION AX502420
VERSION AX502420.1 GI:23384713
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3727 07-AUG-2002;
Aeomica, Inc. (US)
FEATURES
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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
BASE COUNT 9 a 3 c 8 g 5 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 16 CTTCTCTTTT 7
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/db_xref="taxon:9606"
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BASE COUNT
ORIGIN
    9 a _ _ 3 c
    7 g

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 15 CTTCTCTTTT 6

RESULT 41
AX502421/c
LOCUS AX502421 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3728 from Patent EP1229046.
ACCESSION AX502421
VERSION AX502421.1 GI:23384714
KEYWORDS Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Zhan,J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3728 07-AUG-2002;
Aeomica, Inc. (US)
FEATURES
source
    1..25
    /organism="Homo sapiens"
    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
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BASE COUNT
ORIGIN
    9 a _ _ 3 c
    7 g

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 14 CTTCTCTTTT 5

RESULT 42
AX502422/c
LOCUS AX502422 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3729 from Patent EP1229046.
ACCESSION AX502422
VERSION AX502422.1 GI:23384715
KEYWORDS Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Zhan,J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3729 07-AUG-2002;
Aeomica, Inc. (US)
FEATURES
source
    1..25
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    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
6 t
BASE COUNT
ORIGIN
    9 a _ _ 3 c
    7 g

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 11 CTTCTCTTTT 2

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 13 CTTCTCTTTT 4

RESULT 43
AX502423/c
LOCUS AX502423 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3730 from Patent EP1229046.
ACCESSION AX502423
VERSION AX502423.1 GI:23384716
KEYWORDS Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Zhan,J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3730 07-AUG-2002;
Aeomica, Inc. (US)
FEATURES
source
    1..25
    /organism="Homo sapiens"
    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
6 t
BASE COUNT
ORIGIN
    9 a _ _ 3 c
    7 g

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 12 CTTCTCTTTT 3

RESULT 44
AX502424/c
LOCUS AX502424 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3731 from Patent EP1229046.
ACCESSION AX502424
VERSION AX502424.1 GI:23384717
KEYWORDS Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Zhan,J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3731 07-AUG-2002;
Aeomica, Inc. (US)
FEATURES
source
    1..25
    /organism="Homo sapiens"
    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
6 t
BASE COUNT
ORIGIN
    9 a _ _ 4 c
    6 g

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 11 CTTCTCTTTT 2
```

```

RESULT 45
AX502425/c
LOCUS       AX502425                25 bp    DNA
DEFINITION   Sequence 3732 from Patent EP1229046.
ACCESSION    AX502425
VERSION      AX502425.1  GI:23384718
KEYWORDS     Homo sapiens (human)
SOURCE       Homo sapiens
ORGANISM     Homo sapiens
REFERENCE    1
AUTHORS      Zhan, J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 3732 07-AUG-2002;
              Aeomica, Inc. (US)
FEATURES     source
              1..25
              Location/Qualifiers
                /organism="Homo sapiens"
                /mol_type="genomic DNA"
                /db_xref="taxon:9606"
BASE COUNT   9 a      4 c      6 g      6 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 10 CTTCTCTTTT 1

RESULT 46
148977
LOCUS       I48977                25 bp    DNA
DEFINITION   Sequence 4 from patent US 5627054.
ACCESSION    I48977
VERSION      I48977.1  GI:2467440
KEYWORDS     Unknown.
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 25)
AUTHORS      Gillespie, D. deceased.
TITLE        Competitor primer asymmetric polymerase chain reaction
JOURNAL      Patent: US 5627054-A 4 06-MAY-1997;
              Location/Qualifiers
FEATURES     source
              1..25
              /organism="unknown"
BASE COUNT   4 a      5 c      8 t      5 others
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 8 CTTCTCTTTT 17

RESULT 47
ATH525492/c
LOCUS       ATH525492                26 bp    DNA
DEFINITION   Arabidopsis thaliana T-DNA flanking sequence, left border, clone
              057H03.
ACCESSION    AJ525492
VERSION      AJ525492.1  GI:26793728
KEYWORDS     left border; T-DNA flanking sequence.
SOURCE       Arabidopsis thaliana (thale cress)
ORGANISM     Arabidopsis thaliana
              Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
              Eukaryota; Magnoliophyta; eudicotyledons; core eudicots;
              rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE    1
AUTHORS      Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
              Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
              Lepiniec, L., Caboche, M. and Lecharny, A.
TITLE        T-DNA integration into the Arabidopsis genome depends on sequences
              of pre-insertion sites
JOURNAL      EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE      22363535
PUBMED       12446565
REFERENCE    2 (bases 1 to 26)
Direct Submission
Balzerque, S.
Submitted (21-NOV-2002) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
Location/Qualifiers
  1..26
  /organism="Arabidopsis thaliana"
  /mol_type="genomic DNA"
  /cultivar="Wassiliewskija"
  /db_xref="taxon:3702"
  /clone="097H03"
  /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
misc_feature 1..26
              /note="T-DNA flanking sequence"
BASE COUNT   13 a      3 c      4 g      6 t
ORIGIN
Query Match 100.0%; Score 10; DB 8; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 21 CTTCTCTTTT 12

RESULT 48
ATH525503/c
LOCUS       ATH525503                26 bp    DNA
DEFINITION   Arabidopsis thaliana T-DNA flanking sequence, left border, clone
              098B06.
ACCESSION    AJ525503
VERSION      AJ525503.1  GI:26793739
KEYWORDS     left border; T-DNA flanking sequence.
SOURCE       Arabidopsis thaliana (thale cress)
ORGANISM     Arabidopsis thaliana
              Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
              Eukaryota; Magnoliophyta; eudicotyledons; core eudicots;
              rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE    1
AUTHORS      Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
              Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
              Lepiniec, L., Caboche, M. and Lecharny, A.
TITLE        T-DNA integration into the Arabidopsis genome depends on sequences
              of pre-insertion sites
JOURNAL      EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE      22363535
PUBMED       12446565
REFERENCE    2 (bases 1 to 26)

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Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE    1
AUTHORS      Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
              Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
              Lepiniec, L., Caboche, M. and Lecharny, A.
TITLE        T-DNA integration into the Arabidopsis genome depends on sequences
              of pre-insertion sites
JOURNAL      EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE      22363535
PUBMED       12446565
REFERENCE    2 (bases 1 to 26)
Direct Submission
Balzerque, S.
Submitted (21-NOV-2002) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
Location/Qualifiers
  1..26
  /organism="Arabidopsis thaliana"
  /mol_type="genomic DNA"
  /cultivar="Wassiliewskija"
  /db_xref="taxon:3702"
  /clone="097H03"
  /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
misc_feature 1..26
              /note="T-DNA flanking sequence"
BASE COUNT   13 a      3 c      4 g      6 t
ORIGIN
Query Match 100.0%; Score 10; DB 8; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 21 CTTCTCTTTT 12

RESULT 48
ATH525503/c
LOCUS       ATH525503                26 bp    DNA
DEFINITION   Arabidopsis thaliana T-DNA flanking sequence, left border, clone
              098B06.
ACCESSION    AJ525503
VERSION      AJ525503.1  GI:26793739
KEYWORDS     left border; T-DNA flanking sequence.
SOURCE       Arabidopsis thaliana (thale cress)
ORGANISM     Arabidopsis thaliana
              Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
              Eukaryota; Magnoliophyta; eudicotyledons; core eudicots;
              rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE    1
AUTHORS      Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
              Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
              Lepiniec, L., Caboche, M. and Lecharny, A.
TITLE        T-DNA integration into the Arabidopsis genome depends on sequences
              of pre-insertion sites
JOURNAL      EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE      22363535
PUBMED       12446565
REFERENCE    2 (bases 1 to 26)

```

AUTHORS Balzergue,S.  
 TITLE Direct Submission  
 JOURNAL Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue Gaston Cremieux, 91057 Evry cedex, FRANCE

COMMENT PCR was performed on DNA from transformants of Arabidopsis thaliana plants from INRA (Versailles). The DNA fragment(s) resulting from the PCR were directly sequenced from the left or the right border to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed. Information to order the corresponding mutant line and a link to a database providing a graphical display of the insertion site are available at <http://dbsgap.versailles.inra.fr/publiclines/>. This sequence has been generated in the framework of the French plant genomics program "Genoplante" (<http://www.genoplante.com> and <http://genoplante-info.infobiogen.fr>).

FEATURES  
 source  
 1..26  
 /organism="Arabidopsis thaliana"  
 /mol\_type="genomic DNA"  
 /cultivar="Wassilewskija"  
 /db\_xref="taxon:3702"  
 /clone="098B06"  
 /clone\_lib="Arabidopsis thaliana T-DNA insertion lines"  
 misc\_feature  
 1..26  
 /note="T-DNA flanking sequence  
 left border"  
 BASE COUNT 13 a 3 c 4 g 6 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 8; Length 26;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 21 CTTCTCTTTT 12

RESULT 49  
 AR066258/c  
 LOCUS AR066258 27 bp DNA linear PAT 29-SEP-1999  
 DEFINITION Sequence 23 from patent US 5849900.  
 ACCESSION AR066258  
 VERSION AR066258.1 GI:5996474  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unclassified.  
 REFERENCE 1 (bases 1 to 27)  
 AUTHORS Moelling,K.  
 TITLE Inhibition of viruses by antisense oligomers capable of binding to polypurine rich tract of single-stranded RNA or RNA-DNA hybrids  
 JOURNAL Patent: US 5849900-A 23 15-DEC-1998;  
 FEATURES Location/Qualifiers  
 source  
 1..27  
 /organism="unknown"  
 BASE COUNT 7 a 8 c 8 g 4 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 22 CTTCTCTTTT 13

RESULT 50  
 AR191561/c  
 LOCUS AR191561 27 bp DNA linear PAT 20-APR-2002  
 DEFINITION Sequence 7049 from patent US 6346398.  
 ACCESSION AR191561

VERSION AR191561.1 GI:20237526  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 27)  
 AUTHORS Pavco,P., McSwiggen,J., Stinchcomb,D. and Escobedo,J.  
 TITLE Method and reagent for the treatment of diseases or conditions related to levels of vascular endothelial growth factor receptor  
 JOURNAL Patent: US 6346398-A 7049 12-FEB-2002;  
 FEATURES Location/Qualifiers  
 source  
 1..27  
 /organism="unknown"  
 BASE COUNT 12 a 1 c 9 g 4 t 1 others  
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 27 CTTCTCTTTT 18

RESULT 51  
 AX115839/c  
 LOCUS AX115839 27 bp DNA linear PAT 11-MAY-2001  
 DEFINITION Sequence 962 from Patent WO0129262.  
 ACCESSION AX115839  
 VERSION AX115839.1 GI:14032781  
 KEYWORDS  
 SOURCE synthetic construct  
 ORGANISM synthetic construct  
 REFERENCE 1  
 AUTHORS Picoult-Newburg,L. and Pohl,M.  
 TITLE Genotyping reagents, kits and methods of use thereof  
 JOURNAL Patent: WO 0129262-A 962 26-APR-2001;  
 ORCHID Biosciences, Inc. (US)  
 FEATURES Location/Qualifiers  
 source  
 1..27  
 /organism="synthetic construct"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:32630"  
 /note="Primer"  
 BASE COUNT 13 a 5 c 4 g 5 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 21 CTTCTCTTTT 12

RESULT 52  
 AX040823  
 LOCUS AX040823 28 bp DNA linear PAT 23-NOV-2000  
 DEFINITION Sequence 16 from Patent WO0064930.  
 ACCESSION AX040823  
 VERSION AX040823.1 GI:11340462  
 KEYWORDS  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 REFERENCE 1  
 AUTHORS Jay,G.D.  
 TITLE Tribonectins  
 JOURNAL Patent: WO 0064930-A 16 02-NOV-2000;

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FEATURES
  source
    RHODE ISLAND HOSPITAL (US)
    1. .28
    /organism="Homo sapiens"
    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
  BASE COUNT      5 a      7 c      6 g      10 t
  ORIGIN
    Query Match      100.0%; Score 10; DB 6; Length 28;
    Best Local Similarity 100.0%; Pred. No. 1.3e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY      1 CTTCTCTTTT 10
          |||
  Db      12 CTTCTCTTTT 21

  RESULT 53
  AR261655
  LOCUS      29 bp      DNA      PAT 29-JAN-2003
  DEFINITION Sequence 133 from patent US 6322976.
  ACCESSION AR261655
  VERSION AR261655.1 GI:28072733
  KEYWORDS
  SOURCE      Unknown.
  ORGANISM      Unclassified.
  REFERENCE 1 (bases 1 to 29)
  AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
  TITLE Compositions and methods of disease diagnosis and therapy
  JOURNAL Patent: US 6322976-A 133 27-NOV-2001;
  FEATURES
    Location/Qualifiers
    source
      1. .29
      /organism="unknown"
  BASE COUNT      1 a      5 c      10 g      13 t
  ORIGIN
    Query Match      100.0%; Score 10; DB 6; Length 29;
    Best Local Similarity 100.0%; Pred. No. 1.3e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY      1 CTTCTCTTTT 10
          |||
  Db      8 CTTCTCTTTT 17

  RESULT 54
  AR261654
  LOCUS      30 bp      DNA      PAT 29-JAN-2003
  DEFINITION Sequence 132 from patent US 6322976.
  ACCESSION AR261654
  VERSION AR261654.1 GI:28072732
  KEYWORDS
  SOURCE      Unknown.
  ORGANISM      Unclassified.
  REFERENCE 1 (bases 1 to 30)
  AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
  TITLE Compositions and methods of disease diagnosis and therapy
  JOURNAL Patent: US 6322976-A 132 27-NOV-2001;
  FEATURES
    Location/Qualifiers
    source
      1. .30
      /organism="unknown"
  BASE COUNT      1 a      5 c      10 g      14 t
  ORIGIN
    Query Match      100.0%; Score 10; DB 6; Length 30;
    Best Local Similarity 100.0%; Pred. No. 1.3e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY      1 CTTCTCTTTT 10
          |||
  Db      1 CTTCTCTTTT 10
          |||
          8 CTTCTCTTTT 17

  RESULT 55
  AR261656
  LOCUS      30 bp      DNA      PAT 29-JAN-2003
  DEFINITION Sequence 134 from patent US 6322976.
  ACCESSION AR261656
  VERSION AR261656.1 GI:28072734
  KEYWORDS
  SOURCE      Unknown.
  ORGANISM      Unclassified.
  REFERENCE 1 (bases 1 to 30)
  AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
  TITLE Compositions and methods of disease diagnosis and therapy
  JOURNAL Patent: US 6322976-A 134 27-NOV-2001;
  FEATURES
    Location/Qualifiers
    source
      1. .30
      /organism="unknown"
  BASE COUNT      1 a      5 c      10 g      14 t
  ORIGIN
    Query Match      100.0%; Score 10; DB 6; Length 30;
    Best Local Similarity 100.0%; Pred. No. 1.3e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY      1 CTTCTCTTTT 10
          |||
  Db      8 CTTCTCTTTT 17

  RESULT 56
  AR261657
  LOCUS      30 bp      DNA      PAT 29-JAN-2003
  DEFINITION Sequence 135 from patent US 6322976.
  ACCESSION AR261657
  VERSION AR261657.1 GI:28072735
  KEYWORDS
  SOURCE      Unknown.
  ORGANISM      Unclassified.
  REFERENCE 1 (bases 1 to 30)
  AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
  TITLE Compositions and methods of disease diagnosis and therapy
  JOURNAL Patent: US 6322976-A 135 27-NOV-2001;
  FEATURES
    Location/Qualifiers
    source
      1. .30
      /organism="unknown"
  BASE COUNT      1 a      5 c      9 g      15 t
  ORIGIN
    Query Match      100.0%; Score 10; DB 6; Length 30;
    Best Local Similarity 100.0%; Pred. No. 1.3e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY      1 CTTCTCTTTT 10
          |||
  Db      8 CTTCTCTTTT 17

  RESULT 57
  AR120483/c
  LOCUS      33 bp      DNA      PAT 16-MAY-2001
  DEFINITION Sequence 359 from patent US 6159469.
  ACCESSION AR120483
  VERSION AR120483.1 GI:14104059
  KEYWORDS
  SOURCE      Unknown.
  ORGANISM      Unclassified.
  REFERENCE 1 (bases 1 to 33)
  AUTHORS Choi,G.H., Kunsch,C.A., Barash,S.C., Dillon,P.J., Dougherty,B.,
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Fannon,M.R. and Rosen,C.A.
TITLE      Streptococcus pneumoniae antigens and vaccines
JOURNAL    Patent: US 6159469-A 359 12-DEC-2000;
FEATURES   Location/Qualifiers
            source          1..33
            /organism="unknown"
BASE COUNT      14 a      5 c      9 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||
Db      23 CTTCTCTTTT 14

RESULT 58
BD063492/c
LOCUS      BD063492      33 bp      DNA      linear      PAT 27-AUG-2002
DEFINITION Streptococcus pneumoniae antigens and vaccines.
ACCESSION  BD063492
VERSION     BD063492.1 GI:22609095
KEYWORDS   JP 2001505415-A/246.
SOURCE     unidentified
ORGANISM   unclassified.
REFERENCE  1 (bases 1 to 33)
AUTHORS    Kunsch,C.A., Choi,G.H., Johnson,S.L. and Hromockyj,A.
TITLE      Streptococcus pneumoniae antigens and vaccines
JOURNAL    Patent: JP 2001505415-A 246 24-APR-2001;
            HUMAN GENOME SCIENCES INC
COMMENT    PN JP 2001505415-A/246
            PD 24-APR-2001
            PF 30-OCT-1997 JP 1998520667
            PR 31-OCT-1996 US 60/029960
            PI CHARLES A KUNSCH,GIL H CHOI,SYDNOR L JOHNSON,ALEX HROMOCKYJ PC
               C12N15/31,C12N5/18,C12N1/21,C07K14/315,C12Q1/68,A61K39/09, PC
               G01N33/569,
               PC G01N33/68
               CC Strandedness: Double;
               CC Topology: Linear;
               FH Key      Location/Qualifiers.
FEATURES   source          1..33
            /organism="unidentified"
            /mol_type="genomic DNA"
            /db_xref="taxon:32644"
BASE COUNT      14 a      5 c      9 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||
Db      23 CTTCTCTTTT 14

RESULT 59
E36416
LOCUS      E36416      33 bp      DNA      linear      PAT 18-JUN-2001
DEFINITION dna G.
ACCESSION  E36416
VERSION     E36416.1 GI:13022642
KEYWORDS   JP 1999239489-A/3.
SOURCE     Streptococcus pneumoniae
ORGANISM   Bacteria; Firmicutes; Lactobacillales; Streptococcaceae;
            Streptococcus.
REFERENCE  1 (bases 1 to 33)

```

```

Earl,W.M., Deborah,D.J., Ming,H., Richard,R.W. and Ana,R.R.
dna G
TITLE      JP 1999239489-A 3 07-SEP-1999;
JOURNAL    SMITHKLINE BEECHAM CORP
COMMENT    OS Streptococcus pneumoniae
            PN JP 1999239489-A/3
            PD 07-SEP-1999
            PF 21-OCT-1998 JP 1998338366
            PR 21-OCT-1997 US 60/070912
            PI EARL WILLIAM MEI,DEBORAH D JAWASUKI,MING HWANG, PI RICHARD
               ROIDO WOREN,
               PI ANA RISA RENOX
               PC C12N15/09 A61K31/00,A61K31/70,A61K38/00,A61K39/00,A61K48/00,
               PC C07K14/315,
               PC C07K16/12,C12P21/02,G01N33/53//C12P21/08,C12N15/00,A61K37/02
               CC
               FH Key      Location/Qualifiers
               FT source          1..33
               FT /organism="Streptococcus pneumoniae".
FEATURES   source          1..33
            /organism="Streptococcus pneumoniae"
            /mol_type="genomic DNA"
            /db_xref="taxon:1313"
BASE COUNT      8 a      7 c      4 g      14 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||
Db      11 CTTCTCTTTT 20

RESULT 60
AX218433/c
LOCUS      AX218433      38 bp      mRNA      linear      PAT 07-SEP-2001
DEFINITION Sequence 3875 from Patent WO0159103.
ACCESSION  AX218433
VERSION     AX218433.1 GI:15546157
KEYWORDS   .
SOURCE     synthetic construct
            ORGANISM      synthetic construct
                           artificial sequences.
REFERENCE  1
            AUTHORS      Blatt,L., Mcswiggen,J. and Chowrira,B.M.
            TITLE        Method and reagent for the modulation and diagnosis of cd20 and
                           nogo gene expression
            JOURNAL       Patent: WO 0159103-A 3875 16-AUG-2001;
                           RIBOZYME PHARMACEUTICALS, INC. (US) ; Blatt, Lawrence (US) ;
                           McSwiggen, James (US) ; Chowrira, Bharat M. (US)
FEATURES   Location/Qualifiers
            source          1..38
            /organism="synthetic construct"
            /mol_type="mRNA"
            /db_xref="taxon:32630"
            /note="Nucleic Acid"
BASE COUNT      14 a      6 c      14 g      4 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||
Db      38 CTTCTCTTTT 29

RESULT 61
AX220343

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LOCUS       AX220343               38 bp    mRNA          linear          PAT 07-SEP-2001
DEFINITION   Sequence 5785 from Patent WO0159103.
ACCESSION    AX220343
VERSION      AX220343.1   GI:15548067
SOURCE       synthetic construct
            .
ORGANISM     synthetic construct
            .
REFERENCE    1
AUTHORS      Blatt, L., McSwiggen, J. and Chowrira, B.M.
TITLE        Method and reagent for the modulation and diagnosis of cd20 and
            .
JOURNAL      Nogo gene expression
            .
            Patent: WO 0159103-A 5785 16-AUG-2001;
            RIBOZYME PHARMACEUTICALS, INC. (US); Blatt, Lawrence (US) ;
            McSwiggen, James (US) ; Chowrira, Bharat M. (US)
FEATURES     Location/Qualifiers
            1..38
            /organism="synthetic construct"
            /mol_type="mRNA"
            /db_xref="taxon:32630"
            /note="Nucleic Acid"
BASE COUNT   9 a 10 c 9 g 10 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 29 CTTCTCTTTT 38

RESULT 62
AX273375/c
LOCUS       AX273375               38 bp    mRNA          linear          PAT 29-OCT-2001
DEFINITION   Sequence 944 from Patent WO0162911.
ACCESSION    AX273375
VERSION      AX273375.1   GI:16546112
SOURCE       synthetic construct
            .
ORGANISM     synthetic construct
            .
REFERENCE    1
AUTHORS      Jarvis, T., von Carlowitz, I., McSwiggen, J.A., Hamblin, P.A. and
            .
            Ellis, J.H.
TITLE        Method and reagent for the inhibition of grid
            .
JOURNAL      Patent: WO 0162911-A 944 30-AUG-2001;
            RIBOZYME PHARMACEUTICALS, INC. (US) ; GLAXO GROUP LIMITED (GB)
FEATURES     Location/Qualifiers
            1..38
            /organism="synthetic construct"
            /mol_type="mRNA"
            /db_xref="taxon:32630"
            /note="Enzymatic Nucleic Acid"
BASE COUNT   11 a 6 c 13 g 8 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 38 CTTCTCTTTT 29

RESULT 63
BD182426
LOCUS       BD182426               40 bp    DNA          linear          PAT 15-MAY-2003
DEFINITION   Human artificial chromosomes comprising human antibody light chain
            .
            lambda gene, and non-human animals retaining human artificial
            chromosome transmittable to progeny.

```

```

ACCESSION    BD182426
VERSION      BD182426.1   GI:30793344
KEYWORDS     WO 02092812-A/1.
SOURCE       synthetic construct
            .
ORGANISM     synthetic construct
            .
REFERENCE    1 (bases 1 to 40)
AUTHORS      Kuroiwa, Y., Tomizuka, K., Yoshida, H. and Ishida, I.
TITLE        Human artificial chromosomes comprising human antibody light chain
            .
            lambda gene, and non-human animals retaining human artificial
            chromosome transmittable to progeny
            .
            Patent: WO 02092812-A 1 21-NOV-2002;
            KIRIN BREWERY CO LTD, YOSHIMI KUROIWA, KAZUMA TOMIZUKA, HITOSHI
            YOSHIDA, ISAO ISHIDA
COMMENT      OS Artificial Sequence
            .
            PN WO 02092812-A/1
            .
            PF 10-MAY-2002 WO 2002JP004587
            .
            PR 11-MAY-2001 JP 01P 142371
            .
            PI YOSHIMI KUROIWA, KAZUMA TOMIZUKA, HITOSHI YOSHIDA, ISAO ISHIDA PC
            .
            CC C12N15/09,A01K67/027,C07K16/00,C12P21/08
            .
            FT Key Location/Qualifiers
            .
            FT source 1..40
            .
            FT Location/Qualifiers
            .
            FT /organism="Artificial Sequence".
            .
            FT /organism="synthetic construct"
            .
            FT /mol_type="genomic DNA"
            .
            FT /db_xref="taxon:32630"
BASE COUNT   5 a 14 c 6 g 15 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 40;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 21 CTTCTCTTTT 30

RESULT 64
I12092
LOCUS       I12092               42 bp    DNA          linear          PAT 26-JUL-1995
DEFINITION   Sequence 33 from patent US 5420027.
ACCESSION    I12092
VERSION      I12092.1   GI:909590
SOURCE       Unknown.
            .
ORGANISM     Unclassified.
            .
REFERENCE    1 (bases 1 to 42)
AUTHORS      Fisher, C.W., Barnes, H.J. and Estabrook, R.W.
TITLE        Methods and compositions for the expression of biologically active
            .
            fusion proteins comprising a eukaryotic cytochrome P450 fused to a
            reductase in bacteria
            .
            Patent: US 5420027-A 33 30-MAY-1995;
            JOURNAL      Patent: US 5420027-A 33 30-MAY-1995;
            .
            FEATURES     Location/Qualifiers
            .
            source 1..42
            .
            /organism="unknown"
BASE COUNT   5 a 9 c 7 g 21 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 42;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 34

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RESULT 65
AX484623
LOCUS
DEFINITION Sequence 1923 from Patent WO02053728.
ACCESSION
VERSION AX484623.1 GI:22318975
KEYWORDS
SOURCE
ORGANISM Candida albicans
Candida albicans
Bukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
AUTHORS Roemer, T., Jiang, B., Boone, C., Bussey, H. and Ohlsen, K.L.
TITLE Gene disruption methodologies for drug target discovery
JOURNAL Patent: WO 02053728-A 1923 11-JUL-2002;
Elitra Pharmaceuticals, Inc. (US)
FEATURES
source
Location/Qualifiers
1..43
/organism="Candida albicans"
/mol_type="genomic DNA"
/db_xref="taxon:5476" 29 t
BASE COUNT 7 a 5 c 2 g 29 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 43;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 12 CTTCTCTTTT 21
|||||

RESULT 66
AX061875/c
LOCUS
DEFINITION Sequence 8 from Patent WO0078978.
ACCESSION AX061875
VERSION AX061875.1 GI:12539921
KEYWORDS
SOURCE
ORGANISM
synthetic construct
synthetic construct
artificial sequences.
REFERENCE
AUTHORS Miller, B.G., Sloan, J.S., Raymond, C.K. and Vanaja, E.
TITLE Pichia methanolica glycerolaldehyde-3-phosphate dehydrogenase 1
JOURNAL Patent: WO 0078978-A 8 28-DEC-2000;
ZymoGenetics, Inc. (US) ; Miller, Brady G. (US) ; Sloan, James S. (US)
FEATURES
source
Location/Qualifiers
1..45
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/Note="oligonucleotide primer ZC12,565"
BASE COUNT 23 a 8 g 5 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 16 CTTCTCTTTT 7
|||||

RESULT 67
ATH528602/c
LOCUS
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone 168E06.

```

```

ACCESSION AJ528602
VERSION AJ528602.1 GI:26796862
KEYWORDS left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Arabidopsi
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi
1
Brunaud, V., Balzergue, S., Dubreucq, B., Aubourg, S., Samson, P.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepiniec, L., Caboche, M. and Lecharny, A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
PUBMED 12363535
REFERENCE 2 (bases 1 to 45)
12446565
Balzergue, S.
Direct Submission
Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbsgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
Location/Qualifiers
1..45
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassiljewskija"
/db_xref="taxon:3702"
/clone="168E06"
/Note="T-DNA flanking sequence"
misc_feature 1..45
/Note="T-DNA flanking sequence"
left border"
BASE COUNT 16 a 8 c 9 g 12 t
ORIGIN
Query Match 100.0%; Score 10; DB 8; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 10 CTTCTCTTTT 1
|||||

RESULT 68
AR284713
LOCUS
DEFINITION Sequence 765 from patent US 6528260.
ACCESSION AR284713
VERSION AR284713.1 GI:29721617
KEYWORDS
SOURCE
ORGANISM
Unknown.
Unknown.
Unclassified.
REFERENCE 1 (bases 1 to 47)
AUTHORS Blumenfeld, M., Chumakov, I., Bougueleret, L. and Cohen, A.
TITLE Biallelic markers related to genes involved in drug metabolism
JOURNAL Patent: US 6528260-A 765 04-MAR-2003;
Location/Qualifiers
1..47
/organism="unknown"

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BASE COUNT 6 a 11 c 8 g 21 t 1 others  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 6; Length 47;  
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 1 CTTCTCTTTT 10  
 RESULT 69  
 AR288787/c  
 LOCUS AR288787 47 bp DNA PAT 12-JUN-2003  
 DEFINITION Sequence 522 from patent US 6537751.  
 ACCESSION AR288787  
 VERSION AR288787.1 GI:31676071  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.  
 TITLE Biallelic markers for use in constructing a high density  
 disequilibrium map of the human genome  
 JOURNAL Patent: US 6537751-A 522 25-MAR-2003;  
 FEATURES Location/Qualifiers  
 source  
 1..47  
 /organism="unknown"  
 BASE COUNT 16 a 4 c 18 g 8 t 1 others  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 6; Length 47;  
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 15 CTTCTCTTTT 6  
 RESULT 70  
 AR289547  
 LOCUS AR289547 47 bp DNA PAT 12-JUN-2003  
 DEFINITION Sequence 1282 from patent US 6537751.  
 ACCESSION AR289547  
 VERSION AR289547.1 GI:31676831  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.  
 TITLE Biallelic markers for use in constructing a high density  
 disequilibrium map of the human genome  
 JOURNAL Patent: US 6537751-A 1282 25-MAR-2003;  
 FEATURES Location/Qualifiers  
 source  
 1..47  
 /organism="unknown"  
 BASE COUNT 9 a 12 c 10 g 15 t 1 others  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 6; Length 47;  
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 13 CTTCTCTTTT 22  
 RESULT 71

AR289811  
 LOCUS AR289811 47 bp DNA PAT 12-JUN-2003  
 DEFINITION Sequence 1546 from patent US 6537751.  
 ACCESSION AR289811  
 VERSION AR289811.1 GI:31677095  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.  
 TITLE Biallelic markers for use in constructing a high density  
 disequilibrium map of the human genome  
 JOURNAL Patent: US 6537751-A 1546 25-MAR-2003;  
 FEATURES Location/Qualifiers  
 source  
 1..47  
 /organism="unknown"  
 BASE COUNT 5 a 14 c 24 t 2 others  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 6; Length 47;  
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 9 CTTCTCTTTT 18  
 RESULT 72  
 AR290737/c  
 LOCUS AR290737 47 bp DNA PAT 12-JUN-2003  
 DEFINITION Sequence 2472 from patent US 6537751.  
 ACCESSION AR290737  
 VERSION AR290737.1 GI:31678021  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.  
 TITLE Biallelic markers for use in constructing a high density  
 disequilibrium map of the human genome  
 JOURNAL Patent: US 6537751-A 2472 25-MAR-2003;  
 FEATURES Location/Qualifiers  
 source  
 1..47  
 /organism="unknown"  
 BASE COUNT 24 a 6 c 7 g 9 t 1 others  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 6; Length 47;  
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 45 CTTCTCTTTT 36  
 RESULT 73  
 AR291549/c  
 LOCUS AR291549 47 bp DNA PAT 12-JUN-2003  
 DEFINITION Sequence 3284 from patent US 6537751.  
 ACCESSION AR291549  
 VERSION AR291549.1 GI:31678833  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.  
 TITLE Biallelic markers for use in constructing a high density  
 disequilibrium map of the human genome

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JOURNAL Patent: US 6537751-A 3284 25-MAR-2003;
FEATURES
  source
    Location/Qualifiers
      1. .47
        /organism="unknown"
BASE COUNT 19 a 4 c 12 g 11 t 1 others
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 74
AR139658/c
LOCUS AR139658 50 bp DNA linear PAT 16-JUN-2001
DEFINITION Sequence 37 from patent US 6207389.
ACCESSION AR139658
VERSION AR139658.1 GI:14482154
KEYWORDS
SOURCE Unknown.
ORGANISM
REFERENCE
  AUTHORS Unclassified.
  TITLE 1 (bases 1 to 50)
  JOURNAL Dosch,H.Michael.
  FEATURES
    source
      Location/Qualifiers
        1. .50
          /organism="unknown"
BASE COUNT 20 a 8 c 16 g 6 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 42 CTTCTCTTTT 33

RESULT 75
AR158156
LOCUS AR158156 50 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 1484 from Patent WO0140521.
ACCESSION AR158156
VERSION AR158156.1 GI:14539487
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
  AUTHORS Homo sapiens
  TITLE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  JOURNAL Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
  FEATURES
    source
      Location/Qualifiers
        1. .50
          /organism="Homo sapiens"
          /mol_type="genomic DNA"
          /db_xref="taxon:9606"
      misc_feature
        25..26
          /note="Nucleotide deleted between bases 25 and 26"
      misc_feature
        26
          /note="Nucleotide deleted between bases 25 and 26"
BASE COUNT 4 a 9 c 2 g 35 t

JOURNAL Patent: US 6537751-A 3284 25-MAR-2003;
FEATURES
  source
    Location/Qualifiers
      1. .47
        /organism="unknown"
BASE COUNT 19 a 4 c 12 g 11 t 1 others
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 76
AR118141
LOCUS AR118141 51 bp DNA linear PAT 11-MAY-2001
DEFINITION Sequence 3264 from Patent WO0129262.
ACCESSION AR118141
VERSION AR118141.1 GI:14035092
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
  AUTHORS Homo sapiens
  TITLE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  JOURNAL Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
  FEATURES
    source
      Location/Qualifiers
        1. .51
          /organism="Homo sapiens"
          /mol_type="genomic DNA"
          /db_xref="taxon:9606"
BASE COUNT 8 a 13 c 3 g 27 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 20

RESULT 77
AR158155
LOCUS AR158155 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 1483 from Patent WO0140521.
ACCESSION AR158155
VERSION AR158155.1 GI:14539486
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
  AUTHORS Homo sapiens
  TITLE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  JOURNAL Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
  FEATURES
    source
      Location/Qualifiers
        1. .51
          /organism="Homo sapiens"
          /mol_type="genomic DNA"
          /db_xref="taxon:9606"
      misc_feature
        26
          /note="1 of 2 allelic variants (1484 is other entry)"
      misc_feature
        26
          /note="1 of 2 allelic variants (1484 is other entry)"
BASE COUNT 4 a 9 c 2 g 36 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 51;

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Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 27 CTTCTCTTTT 36

RESULT 78
AX160381
LOCUS AX160381 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3709 from Patent WO0140521.
ACCESSION AX160381
VERSION AX160381.1 GI:14541712
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0140521-A 3709 07-JUN-2001;
Curagen Corporation (US)
FEATURES
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/organism="Homo sapiens"
/mol_type="genomic DNA"
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misc_feature 26
/notes="1 of 2 allelic variants (3710 is other entry)
Accession number CG43917418"
BASE COUNT 7 a 15 c 10 g 19 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 40 CTTCTCTTTT 49

RESULT 79
AX160382
LOCUS AX160382 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3710 from Patent WO0140521.
ACCESSION AX160382
VERSION AX160382.1 GI:14541713
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0140521-A 3710 07-JUN-2001;
Curagen Corporation (US)
FEATURES
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/db_xref="taxon:9606"
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/notes="2 of 2 allelic variants (3709 is other entry)
Accession number CG43917418"
BASE COUNT 7 a 16 c 10 g 18 t
ORIGIN

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Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 40 CTTCTCTTTT 49

RESULT 80
AX160383
LOCUS AX160383 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3711 from Patent WO0140521.
ACCESSION AX160383
VERSION AX160383.1 GI:14541714
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0140521-A 3711 07-JUN-2001;
Curagen Corporation (US)
FEATURES
source
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/db_xref="taxon:9606"
misc_feature 26
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Accession number CG43917418"
BASE COUNT 6 a 12 c 10 g 23 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 34

RESULT 81
AX165562
LOCUS AX165562 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 757 from Patent WO0138586.
ACCESSION AX165562
VERSION AX165562.1 GI:14546391
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0138586-A 757 31-MAY-2001;
Curagen Corporation (US)
FEATURES
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/db_xref="taxon:9606"
variation 26
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BASE COUNT 12 a 12 c 7 g 20 t
ORIGIN

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Query Match      100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11

RESULT 82
ATH521150
LOCUS ATH521150 51 bp DNA linear PLN 29-MAR-2003
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone 260A09.
ACCESSION AJ521150 GI:26789386
VERSION AJ521150.1
KEYWORDS left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1
Brunaud,V., Balzerque,S., Dubreucq,B., Aubourg,S., Samson,F.,
Chauvin,S., Bechtold,N., Cruaud,C., DeRose,R., Pelletier,G.,
Lepiniec,L., Caboche,M. and Lecharny,A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
23363535
12446565
REFERENCE 2 (bases 1 to 51)
AUTHORS Balzerque,S.
TITLE Direct Submission
JOURNAL Submitted (21-NOV-2002) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
Location/Qualifiers
1..51
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassilewskija"
/db_xref="taxon:3702"
/clone="260A09"
/misc_feature 1..51
/note="T-DNA flanking sequence
left border"
BASE COUNT 15 a 9 c 2 g 25 t
ORIGIN
Query Match      100.0%; Score 10; DB 8; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 4 CTTCTCTTTT 13

RESULT 83
AR098682/c
LOCUS AR098682 53 bp DNA linear PAT 14-FEB-2001
DEFINITION Sequence 40 from patent US 6077668.
ACCESSION AR098682
VERSION AR098682.1 GI:12808448
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 53)
AUTHORS Kool,E.T.
TITLE Highly sensitive multimeric nucleic acid probes
JOURNAL Patent: US 6077668-A 40 20-JUN-2000;
FEATURES Location/Qualifiers
1..53
/organism="unknown"
BASE COUNT 20 a 10 c 15 g 8 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 84
AR098683
LOCUS AR098683 53 bp DNA linear PAT 14-FEB-2001
DEFINITION Sequence 41 from patent US 6077668.
ACCESSION AR098683
VERSION AR098683.1 GI:12808449
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 53)
AUTHORS Kool,E.T.
TITLE Highly sensitive multimeric nucleic acid probes
JOURNAL Patent: US 6077668-A 41 20-JUN-2000;
FEATURES Location/Qualifiers
1..53
/organism="unknown"
BASE COUNT 8 a 15 c 10 g 20 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 38 CTTCTCTTTT 47

RESULT 85
AR204756/c
LOCUS AR204756 53 bp DNA linear PAT 20-JUN-2002
DEFINITION Sequence 40 from patent US 6388802.
ACCESSION AR204756
VERSION AR204756.1 GI:21502164
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 53)
AUTHORS Kool,E.T.
TITLE Circular DNA vectors for synthesis of RNA and DNA
JOURNAL Patent: US 6388802-A 40 09-APR-2002;
FEATURES Location/Qualifiers
1..53
/organism="unknown"
source

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BASE COUNT      20 a      10 c      15 g      8 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      |||||
25 CTTCTCTTTT 16

RESULT 86
AR204757
LOCUS      AR204757      53 bp      DNA      linear      PAT 20-JUN-2002
DEFINITION      Sequence 41 from patent US 6368802.
ACCESSION      AR204757
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unclassified.
REFERENCE      1 (bases 1 to 53)
AUTHORS      Kool,E.T.
TITLE      Circular DNA vectors for synthesis of RNA and DNA
JOURNAL      Patent: US 6368802-A 41 09-APR-2002;
FEATURES
source      1..53
/organism="unknown"
BASE COUNT      8 a      15 c      10 g      20 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      |||||
38 CTTCTCTTTT 47

RESULT 87
AR134108/c
LOCUS      AR134108      54 bp      DNA      linear      PAT 16-MAY-2001
DEFINITION      Sequence 2533 from patent US 6194150.
ACCESSION      AR134108
VERSION      AR134108.1 GI:14123013
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unclassified.
REFERENCE      1 (bases 1 to 54)
AUTHORS      Stinchcomb,D.T., Jarvis,T. and McSwiggen,J.
TITLE      Nucleic acid based inhibition of CD40
JOURNAL      Patent: US 6194150-A 2533 27-FEB-2001;
FEATURES
source      1..54
/organism="unknown"
BASE COUNT      20 a      12 c      13 g      9 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      |||||
13 CTTCTCTTTT 4

RESULT 88
AR134285/c
LOCUS      AR134285      54 bp      DNA      linear      PAT 16-MAY-2001
DEFINITION      Sequence 2710 from patent US 6194150.
ACCESSION      AR134285
VERSION      AR134285.1 GI:14123190
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unclassified.
REFERENCE      1 (bases 1 to 54)
AUTHORS      Stinchcomb,D.T., Jarvis,T. and McSwiggen,J.
TITLE      Nucleic acid based inhibition of CD40
JOURNAL      Patent: US 6194150-A 2710 27-FEB-2001;
FEATURES
source      1..54
/organism="unknown"
BASE COUNT      21 a      11 c      12 g      10 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      |||||
13 CTTCTCTTTT 4

RESULT 89
ATH505724
LOCUS      ATH505724      57 bp      mRNA      linear      PLN 30-APR-2003
DEFINITION      Arabidopsis thaliana partial mitochondrial small non-messenger RNA,
clone Ath-647, 5' end incomplete.
ACCESSION      AJ505724
VERSION      AJ505724.1 GI:22293620
KEYWORDS      small non-messenger RNA; smRNA.
SOURCE      Mitochondrion Arabidopsis thaliana (thale cress)
ORGANISM      Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE      1
AUTHORS      Marker,C., Zemann,A., Terhorst,T., Kiefmann,M., Kastenmayer,J.P.,
Green,P., Bachelierie,J.P., Brosius,J. and Huttenhofer,A.
TITLE      Experimental RNomics. Identification of 140 Candidates for Small
Non-Messenger RNAs in the Plant Arabidopsis thaliana
JOURNAL      Curr. Biol. 12 (23), 2002-2013 (2002)
MEDLINE      22365595
PUBMED      12477388
REFERENCE      2 (bases 1 to 57)
AUTHORS      Huttenhofer,A.
TITLE      Direct Submission
JOURNAL      Submitted (22-JUL-2002) Huttenhofer A., University of Muenster,
Institute for Experimental Pathology, Von-Esmarch-Str. 56, 48149
Muenster, GERMANY
FEATURES
source      1..57
/organism="Arabidopsis thaliana"
/organelle="mitochondrion"
/mol_type="mRNA"
/db_xref="taxon:3702"
/clone="Ath-647"
misc_RNA
/note="small non-messenger RNA, smRNA"
BASE COUNT      6 a      10 c      8 g      33 t
ORIGIN
Query Match      100.0%; Score 10; DB 8; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      |||||
45 CTTCTCTTTT 54

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RESULT 90
ATH527067/c
LOCUS
DEFINITION
  ATH527067          58 bp      DNA      linear      PLN 29-MAR-2003
  Arabidopsis thaliana T-DNA flanking sequence, left border, clone
  131H07.
ACCESSION
  AJ527067          GI:26795327
VERSION
  AJ527067.1
KEYWORDS
  left border; T-DNA flanking sequence.
SOURCE
  Arabidopsis thaliana (thale cress)
ORGANISM
  Arabidopsis thaliana
  Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
  Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
  rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
  1
  Brunaud,V., Balzergue,S., Dubreucq,B., Aubourg,S., Samson,F.,
  Chauvin,S., Bechtold,N., Cruaud,C., DeRose,R., Pelletier,G.,
  Lepiniec,L., Caboche,M. and Lecharny,A.
  T-DNA integration into the Arabidopsis genome depends on sequences
  of pre-insertion sites
  EMBO Rep. 3 (12), 1152-1157 (2002)
JOURNAL
  22363535
MEDLINE
  12446565
PUBMED
  2
  (bases 1 to 58)
REFERENCE
  Balzergue,S.
  Direct Submission
  Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue
  Gaston Cremieux, 91057 Evry cedex, FRANCE
  PCR was performed on DNA from transformants of Arabidopsis thaliana
  plants from INRA (Versailles). The DNA fragment (s) resulting from
  the PCR were directly sequenced from the left or the right border
  to determine the genomic sequence flanking the insertion. T-DNA
  derived sequences were removed. Information to order the
  corresponding mutant line and a link to a database providing a
  graphical display of the insertion site are available at
  http://dbsgap.versailles.inra.fr/publiclines/. This sequence has
  been generated in the framework of the French plant genomics
  program 'Genoplante' (http://www.genoplante.com and
  http://genoplante-info.infobiogen.fr).
FEATURES
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  /cultivar="Wassilewskija"
  /db_xref="taxon:3702"
  /clone="131H07"
  /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
  misc_feature
  1..58
  /note="T-DNA flanking sequence
  left border"
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  Query Match          100.0%; Score 10; DB 8; Length 58;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 47 CTTCTCTTTT 38

RESULT 91
AX270701
LOCUS
DEFINITION
  Sequence 1332 from Patent WO0164876.
ACCESSION
  AX270701
VERSION
  AX270701.1
KEYWORDS
  Homo sapiens (human)
SOURCE
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

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REFERENCE
  1
  Stefansson,H., Steinhorsdottir,V. and Gulcher,J.R.
  Human schizophrenia gene
  Patent: WO 0164876-A 1332 07-SEP-2001;
  Decode Genetics EHF. (IS)
FEATURES
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  1..61
  /organism="Homo sapiens"
  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
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  2 a      23 c      0 g      35 t      1 others
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  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 92
AX272232
LOCUS
  AX272232
DEFINITION
  Sequence 1332 from Patent WO0164877.
ACCESSION
  AX272232
VERSION
  AX272232.1
KEYWORDS
  Homo sapiens (human)
SOURCE
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
  1
  Stefansson,H., Steinhorsdottir,V. and Gulcher,J.R.
  Human schizophrenia gene
  Patent: WO 0164877-A 1332 07-SEP-2001;
  Decode Genetics EHF. (IS)
FEATURES
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  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 93
AX482835
LOCUS
  AX482835
DEFINITION
  Sequence 135 from Patent WO02053728.
ACCESSION
  AX482835
VERSION
  AX482835.1
KEYWORDS
  Candida albicans
SOURCE
  Candida albicans
  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
  Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
  1
  Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
  Gene disruption methodologies for drug target discovery
  Patent: WO 02053728-A 135 11-JUL-2002;
  Elitra Pharmaceuticals, Inc. (US)
FEATURES
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  1..65
  /organism="Homo sapiens"
  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 94
AX482835
LOCUS
  AX482835
DEFINITION
  Sequence 135 from Patent WO02053728.
ACCESSION
  AX482835
VERSION
  AX482835.1
KEYWORDS
  Candida albicans
SOURCE
  Candida albicans
  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
  Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
  1
  Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
  Gene disruption methodologies for drug target discovery
  Patent: WO 02053728-A 135 11-JUL-2002;
  Elitra Pharmaceuticals, Inc. (US)
FEATURES
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  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
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ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 95
AX482835
LOCUS
  AX482835
DEFINITION
  Sequence 135 from Patent WO02053728.
ACCESSION
  AX482835
VERSION
  AX482835.1
KEYWORDS
  Candida albicans
SOURCE
  Candida albicans
  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
  Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
  1
  Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
  Gene disruption methodologies for drug target discovery
  Patent: WO 02053728-A 135 11-JUL-2002;
  Elitra Pharmaceuticals, Inc. (US)
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  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
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  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

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/organism="Candida albicans"
/mol_type="genomic DNA"
/db_xref="taxon:5476"
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BASE COUNT      5 a      22 c      2 g
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 10; DB 6; Length 65;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11

RESULT 94
AX482852
LOCUS      65 bp      DNA      linear      PAT 16-AUG-2002
DEFINITION Sequence 152 from Patent WO202053728.
ACCESSION AX482852
VERSION    AX482852.1 GI:22317272
KEYWORDS
SOURCE     Candida albicans
ORGANISM   Candida albicans
REFERENCE  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
            Saccharomycetales; mitosporic Saccharomycetales; Candida.
AUTHORS    Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
TITLE      Gene disruption methodologies for drug target discovery
JOURNAL    Patent: WO 02053728-A 152 11-JUL-2002;
            Elitra Pharmaceuticals, Inc. (US)
FEATURES   Location/Qualifiers
            source
              1..65
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              /db_xref="taxon:5476"
BASE COUNT  17 a      16 c      5 g      27 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 10; DB 6; Length 65;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 15 CTTCTCTTTT 24

RESULT 95
AX485490
LOCUS      65 bp      DNA      linear      PAT 16-AUG-2002
DEFINITION Sequence 2790 from Patent WO202053728.
ACCESSION AX485490
VERSION    AX485490.1 GI:22319774
KEYWORDS
SOURCE     Candida albicans
ORGANISM   Candida albicans
REFERENCE  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
            Saccharomycetales; mitosporic Saccharomycetales; Candida.
AUTHORS    Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
TITLE      Gene disruption methodologies for drug target discovery
JOURNAL    Patent: WO 02053728-A 2790 11-JUL-2002;
            Elitra Pharmaceuticals, Inc. (US)
FEATURES   Location/Qualifiers
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QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 34

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LOCUS      70 bp      DNA      linear      PAT 21-JAN-2000
DEFINITION Sequence 1 from Patent WO9900504.
ACCESSION A81696
VERSION    A81696.1 GI:6731831
KEYWORDS
SOURCE     unidentified
ORGANISM   unidentified
REFERENCE  1 (bases 1 to 70)
AUTHORS    Sleep,D.
TITLE      IMPROVED PROTEIN EXPRESSION STRAINS
JOURNAL    Patent: WO 9900504-A 1 07-JAN-1999;
            DELTA BIOTECHNOLOGY LTD (GB); SLEEP DARRELL (GB)
FEATURES   Location/Qualifiers
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Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 30 CTTCTCTTTT 39

RESULT 97
AR207788
LOCUS      70 bp      DNA      linear      PAT 20-JUN-2002
DEFINITION Sequence 1 from patent US 6379924.
ACCESSION AR207788
VERSION    AR207788.1 GI:21507632
KEYWORDS
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 70)
AUTHORS    Sleep,D.
TITLE      Protein expression strains
JOURNAL    Patent: US 6379924-A 1 30-APR-2002;
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Db 30 CTTCTCTTTT 39

RESULT 98
162432
LOCUS      71 bp      DNA      linear      PAT 07-OCT-1997
DEFINITION Sequence 5 from patent US 5659122.

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ACCESSION I62432
VERSION I62432.1 GI:2480380
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 71)
AUTHORS Austin,G.Douglas.
TITLE Enhanced expression in plants using non-translated leader sequences
JOURNAL Patent: US 5659122-A 5 19-AUG-1997;
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BASE COUNT 28 a 17 c 5 g 21 t
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Db 43 CTTCTCTTTT 52

RESULT 99
AH006998S06
LOCUS AH006998S06 71 bp DNA linear PRI 27-SEP-2002
DEFINITION Homo sapiens mitochondrial short-chain L-3-hydroxyacyl-CoA
dehydrogenase (HADHSC) gene, nuclear gene encoding mitochondrial
protein, 5' end of intron 3.
ACCESSION AF026858
VERSION AF026858.1 GI:3882441
KEYWORDS
SEGMENT
SOURCE 6 of 15
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 71)
AUTHORS Vredendaal,P.J., van den Berg,I.E., Stroobants,A.K., van der
A,D.L., Malingre,H.E. and Berger,R.
TITLE Structural organization of the human short-chain
L-3-hydroxyacyl-CoA dehydrogenase gene
JOURNAL Mamm. Genome 9 (9), 763-768 (1998)
MEDLINE 9716664
PUBMED
REFERENCE 2 (bases 1 to 71)
AUTHORS Vredendaal,P.J.C.M., van den Berg,I.E.T., Stroobants,A.K., van der
A,D.L., Malingre,H.E.M. and Berger,R.
TITLE Direct Submision
JOURNAL Submitted (25-SEP-1997) Department of Metabolic Diseases,
Wilhelmina Children's Hospital, Nieuwe Gracht 137, Utrecht 3512 LK,
The Netherlands
FEATURES
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Db 21 CTTCTCTTTT 30

RESULT 100
BD055583/c
LOCUS BD055583 73 bp DNA linear PAT 27-AUG-2002
DEFINITION Sequence tag and encoded human protein.
ACCESSION BD055583
VERSION BD055583.1 GI:22601189
KEYWORDS JP 2001269182-A/31829.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 73)
AUTHORS Edwards,J.B.D.M., Duclair,E. and Jordan,J.Y.
TITLE Sequence tag and encoded human protein
JOURNAL Patent: JP 2001269182-A 31829 02-OCT-2001;
GENSET
COMMENT OS Homo sapiens (human)
PN JP 2001269182-A/31829
PD 02-OCT-2001
PF 24-FEB-2000 JP 2000118773
PR 26-FEB-1999 US 60/122487
PI JEAN BAPTISTE DUMAS MILNE EDWARDS,EIMERIC DUCLAIR,JEAN YVES
PI JORDAN
PC C12N15/09,C07K14/435,C07K16/18,C12N1/15,C12N1/19,C12N1/21, PC
C12N5/10,
PC C12P21/02,C12P21/08,C12Q1/68//G06F17/30,C12N15/00,C12N5/00, PC
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    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 29 CTTCTCTTTT 20

Search completed: October 28, 2003, 17:43:55
Job time : 1553 secs

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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:10:20 : Search time 272 Seconds  
(without alignments)  
99.244 Million cell updates/sec

Title: US-09-335-032-71

Perfect score: 10

Sequence: 1 cttctctttt 10

Scoring table: OLIGO\_NUC

Gapop 60.0 , Gapext 60.0

Searched: 2552756 seqs, 1349719017 residues

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Minimum DB seq length: 0

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Post-processing: Listing first 500 summaries

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Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

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2	10	100.0	10	AAF33332	Yeast NORF gene SA
3	10	100.0	10	AAF33321	Yeast NORF gene SA
4	10	100.0	10	AAF33324	Yeast NORF gene SA
5	10	100.0	10	AAF34233	Yeast NORF gene SA
6	10	100.0	12	AAF34233	Triple helix formi
7	10	100.0	12	AAV14794	Triple helix third
8	10	100.0	12	AAH67817	Oligonucleotide pr

9	10	100.0	12	23	ABI80052	Oligonucleotide pr
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c 13	10	100.0	13	23	ABC53773	Oligonucleotide SE
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C 83	10	100.0	25	24	ABS75583	Human PAPP-Ea asso	C 156	10	100.0	60	10	AA91187	Alpha-factor leader
C 84	10	100.0	25	24	ABS75584	Human PAPP-Ea asso	C 157	10	100.0	60	24	ABN45634	Human spliced tran
C 85	10	100.0	25	24	ABS75585	Human PAPP-Ea asso	C 158	10	100.0	60	24	ABN59213	Human spliced tran
C 86	10	100.0	25	24	ABS75586	Human PAPP-Ea asso	C 159	10	100.0	61	22	AAK36533	Human neuroregulin g
C 87	10	100.0	25	24	ABS75587	Human PAPP-Ea asso	C 160	10	100.0	61	22	AAK39026	Human neuroregulin g
C 88	10	100.0	25	24	ABS75588	Human PAPP-Ea asso	C 161	10	100.0	61	24	ABT01303	Human neuroregulin-1
C 89	10	100.0	25	24	ABS75589	Human PAPP-Ea asso	C 162	10	100.0	61	24	ABT02796	Human neuroregulin-1
C 90	10	100.0	25	24	ABS75590	Human PAPP-Ea asso	C 163	10	100.0	62	20	AAH86211	Human neuroregulin-1
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C 93	10	100.0	27	18	AAQ45398	Mouse flt-1 VSGF r	C 166	10	100.0	63	25	AAH55167	Bovine EST associa
C 94	10	100.0	27	22	AAH2997	Shrimp white spot	C 167	10	100.0	65	24	ABZ26256	Candida essential
C 95	10	100.0	27	22	AAH2997	SNP specific lower	C 167	10	100.0	65	24	ABZ26273	Candida essential
C 96	10	100.0	28	22	AAH38166	Human MSF exon 6 R	C 168	10	100.0	65	24	ABZ28775	Candida gene relat
C 97	10	100.0	29	21	AAAF05757	Hammerhead ribozym	C 169	10	100.0	65	24	ABN54867	Mouse spliced tran
C 98	10	100.0	29	21	AAAF05757	Hammerhead ribozym	C 170	10	100.0	66	20	AAH55313	Mouse spliced tran
C 99	10	100.0	30	21	AAAF06662	Human CD36 polymor	C 171	10	100.0	66	20	AAH86210	Human single nucle
C 100	10	100.0	30	21	AAAF06662	Human CD36 polymor	C 172	10	100.0	66	21	AAH44379	PCR primer for a f
C 101	10	100.0	30	21	AAAF06665	Human CD36 polymor	C 173	10	100.0	67	21	AAAF4896	Sense strand nucle
C 102	10	100.0	31	16	AAAT05588	Human CD36 polymor	C 174	10	100.0	70	20	AAQ99879	S. cerevisiae UBC
C 103	10	100.0	31	16	AAAT05588	Mouse brain p69 cD	C 175	10	100.0	71	15	AAQ78295	Heat shock protein
C 104	10	100.0	31	25	AAAT05588	Human genomic DNA	C 176	10	100.0	71	16	AAQ81861	Sense strand of th
C 105	10	100.0	31	25	AAAT05588	Human genomic DNA	C 177	10	100.0	71	21	AAZ46993	Soybean HSP17.9 le
C 106	10	100.0	32	15	AAAT07410	Human WTSP10 PCR p	C 178	10	100.0	73	21	AAAC31838	Human secreted pro
C 107	10	100.0	33	19	AAV39876	Primer Vbeta14/Eco	C 179	10	100.0	75	16	AAQ81862	Human secreted pro
C 108	10	100.0	33	24	ABQ35508	Streptococcus pneu	C 180	10	100.0	75	16	AAZ46994	Antisense strand o
C 109	10	100.0	36	23	ABQ85037	PCR primer used to	C 181	10	100.0	76	21	AAAC15151	Soybean HSP17.9 le
C 110	10	100.0	36	23	ABQ85037	Streptococcus pneu	C 182	10	100.0	76	24	AAH831918	Human secreted pro
C 111	10	100.0	38	23	ABK03875	Human WTSP10 PCR p	C 183	10	100.0	78	15	AAQ78302	Soybean GmHSP17.9
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C 113	10	100.0	40	14	AAQ04864	Human NOGO Zinzyme	C 185	10	100.0	83	21	AAAC31898	Soybean HSP17.9-B
C 114	10	100.0	40	21	AAQ36817	VAC-beta gene prim	C 186	10	100.0	83	24	ABN60962	Human secreted pro
C 115	10	100.0	40	25	ABZ70085	Forward PCR primer	C 187	10	100.0	86	22	AAAS23462	Human cancer relat
C 116	10	100.0	41	24	ABK30429	Human shearing fac	C 188	10	100.0	87	22	AAK33769	C. albicans essent
C 117	10	100.0	41	24	ABK30430	Human shearing fac	C 189	10	100.0	87	22	AAK49863	Human brain expres
C 118	10	100.0	42	13	AAQ27383	Bovine l7-alpha-hy	C 190	10	100.0	87	22	AAI26968	Human bone marrow
C 119	10	100.0	42	15	AAQ55121	Human liver P450 2	C 191	10	100.0	87	22	AAI55783	Probe #16901 for g
C 120	10	100.0	42	20	AAK89210	Primer for constru	C 192	10	100.0	87	23	ABSA49502	Probe #24469 used
C 121	10	100.0	43	24	AAH84248	Human cell death p	C 193	10	100.0	87	22	AAI53826	Human liver single
C 122	10	100.0	45	22	AAAD27976	Candida essential	C 194	10	100.0	89	22	ABSA23361	Human genome-deriv
C 123	10	100.0	47	20	AAK03631	Pichia methanolica	C 195	10	100.0	89	23	ABSA47713	Probe #22512 used
C 124	10	100.0	47	21	AAZ66175	Actinia equina L.	C 196	10	100.0	90	24	ABK36650	Human liver single
C 125	10	100.0	47	21	AAZ66175	Human map-related	C 197	10	100.0	90	24	ABK36650	Human DNA encoding
C 126	10	100.0	47	21	AAZ67199	Human map-related	C 198	10	100.0	93	22	ABAT1937	Human brain expres
C 127	10	100.0	47	21	AAZ68125	Human map-related	C 199	10	100.0	93	22	AAK20329	Human foetal liver
C 128	10	100.0	47	21	AAZ68930	Human map-related	C 200	10	100.0	93	22	AAK46413	Human brain expres
C 129	10	100.0	48	22	AAH84247	Human map-related	C 201	10	100.0	93	22	AAK46413	Human bone marrow
C 130	10	100.0	50	21	AAAF77160	Human cell death p	C 202	10	100.0	93	22	ABSA46139	Probe #20984 used
C 131	10	100.0	50	22	AAI74543	Human clone cg4394	C 203	10	100.0	93	23	ABSA46139	Human liver single
C 132	10	100.0	50	22	AAI74543	Human clone cg4394	C 204	10	100.0	93	24	ABSA20735	Human liver single
C 133	10	100.0	50	22	AAI74543	Mouse silent SNP c	C 205	10	100.0	93	25	ABZ78911	Human genome-deriv
C 134	10	100.0	50	24	ABZ02704	Human brain p69 5'	C 206	10	100.0	93	25	ABZ78911	Tumour suppression
C 135	10	100.0	50	24	ABZ04201	Human leukocyte ge	C 207	10	100.0	95	16	AAI23132	Human oligonucleot
C 136	10	100.0	50	24	ABZ04416	Human leukocyte ge	C 208	10	100.0	96	22	AAH84246	Human gene signatu
C 137	10	100.0	50	24	ABZ06159	Human leukocyte ge	C 209	10	100.0	97	15	AAI19418	Human cell death p
C 138	10	100.0	50	24	ABZ06269	Human leukocyte ge	C 210	10	100.0	97	22	ABAT5204	Human gene signatu
C 139	10	100.0	51	21	AAAF77161	Human leukocyte ge	C 211	10	100.0	97	22	ABAT5204	Human foetal liver
C 140	10	100.0	51	22	AAAL27570	Human clone cg4394	C 212	10	100.0	97	22	AAK33739	Probe #18333 for g
C 141	10	100.0	51	22	AAAL30883	Human SNP oligonuc	C 213	10	100.0	97	22	AAK33739	Human brain expres
C 142	10	100.0	51	22	AAAL32705	Human SNP oligonuc	C 214	10	100.0	97	22	AAI26943	Human bone marrow
C 143	10	100.0	51	22	AAAL33361	Human SNP oligonuc	C 215	10	100.0	97	22	AAI26943	Probe #16876 for g
C 144	10	100.0	51	22	AAI74542	Human silent SNP c	C 216	10	100.0	97	23	ABSA49475	Probe #24439 used
C 145	10	100.0	51	22	AAI76768	Human silent SNP c	C 217	10	100.0	97	24	ABSA49475	Human liver single
C 146	10	100.0	51	22	AAI76769	Human silent SNP c	C 218	10	100.0	99	21	AAAC14676	Human genome-deriv
C 147	10	100.0	51	22	AAI76770	Human silent SNP c	C 219	10	100.0	99	21	AAAC14676	Human secreted pro
C 148	10	100.0	51	22	AAH40468	Human silent SNP c	C 220	10	100.0	102	21	AAAT0384	Human secreted pro
C 149	10	100.0	51	23	ABL00766	Human SNP flanking	C 221	10	100.0	102	21	ABAT5552	5' oligonucleotide
C 150	10	100.0	53	19	AAV59245	Human amino acid c	C 222	10	100.0	102	22	ABAT5552	Human foetal liver
C 151	10	100.0	53	19	AAV59246	DNA 53mer circle s	C 223	10	100.0	102	22	ABAT5552	Human foetal liver
C 152	10	100.0	53	19	AAV12931	Stem loop RNA mult	C 224	10	100.0	102	22	AAK39332	Probe #18631 for g
C 153	10	100.0	53	19	AAV12932	Oligonucleotide SE	C 225	10	100.0	102	22	AAI27265	Human brain expres
C 154	10	100.0	53	20	AAK30035	Oligonucleotide SE	C 226	10	100.0	102	22	AAI27265	Human bone marrow
C 155	10	100.0	54	17	AAK67061	Mouse B7 hairpin r	C 227	10	100.0	102	23	ABSA49816	Human liver single



C 374	10	100.0	10	100.0	10	100.0	218	22	AAI21506	Probe #11439 for g
C 375	10	100.0	10	100.0	10	100.0	218	22	AAI46798	Probe #15484 used
C 376	10	100.0	10	100.0	10	100.0	218	22	AAI07202	Probe #7193 used t
C 377	10	100.0	10	100.0	10	100.0	218	22	ABSA0316	Human liver single
C 378	10	100.0	10	100.0	10	100.0	218	23	ABSA0316	Human liver single
C 379	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 380	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 381	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 382	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 383	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 384	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 385	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 386	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 387	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 388	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 389	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 390	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 391	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 392	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 393	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 394	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 395	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 396	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 397	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 398	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 399	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 400	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 401	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 402	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 403	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 404	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 405	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 406	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 407	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 408	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 409	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 410	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 411	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 412	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 413	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 414	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 415	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 416	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 417	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 418	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 419	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 420	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 421	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 422	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 423	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 424	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 425	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 426	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 427	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 428	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 429	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 430	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 431	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 432	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 433	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 434	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 435	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 436	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 437	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 438	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 439	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 440	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 441	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 442	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 443	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 444	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 445	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv
C 446	10	100.0	10	100.0	10	100.0	218	24	ABSA14697	Human genome-deriv

## ALIGNMENTS

## RESULT 1

AAV50115  
 XX AAV50115 standard; DNA; 10 BP.  
 XX  
 AC AAV50115;  
 XX  
 DT 21-OCT-1998 (first entry)  
 XX Yeast tag for NORF gene locus NORF5.  
 DE  
 DE Yeast: Saccharomyces cerevisiae; transcriptome; cell cycle;  
 KW regulation; eukaryotic cell; antifungal; SAGE tag; gene expression;  
 KW serial analysis of gene expression; probe; ss.  
 XX

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OS Saccharomyces cerevisiae.
OS Synthetic.
XX
XX
PN WO9832847-A2.
XX
XX
PD 30-JUL-1998.
XX
XX
PF PF
XX
XX 22-JAN-1998; 98WO-US01216.
XX
XX 23-JAN-1997; 97US-0035917.
XX
XX (UYJO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX Kinzler KW, Velculescu VE, Vogelstein B;
XX
XX WPI; 1998-427943/36.
XX
XX Yeast transcriptome - useful for modulating eukaryotic cell, for
PT screening antifungal agents, and for identifying genes in cell cycle
PT progression
XX
XX Claim 1; Page 23; 44pp; English.
XX
XX Yeast transcriptome is encoded by a DNA molecule comprising a yeast
CC gene involved in cell cycle progression selected from the group of
CC nonannotated ORF (NORF) genes. SAGE (serial analysis of gene expression)
CC tags for highly expressed genes and NORF genes are given in AAV50051 to
CC AAV50345. The present invention describes: (1) a method of using yeast
CC genes to modulate the cell cycle which comprises administering to a cell
CC an isolated DNA molecule comprising a yeast gene which is involved in
CC cell cycle progression selected from differentially expressed genes
CC (SAGE tags given in AAV50051 to AAV50345); (2) a method for screening
CC candidate antifungal drugs which comprises contacting a test substance
CC with a yeast cell and monitoring expression of a yeast gene which is
CC involved in cell cycle progression; (3) a method of identifying human
CC genes which are involved in cell cycle progression which comprises
CC hybridizing a probe comprising at least 10 contiguous nucleotides of a
CC yeast gene which is differentially expressed between at least 2 phases
CC selected from the log phase, the S phase and the G2/M phase; and (4) a
CC probe for ascertaining the phase in the cell cycle, where the probe
CC comprises at least 14 contiguous nucleotides of a NORF gene (SAGE tags
CC given in AAV50051 to AAV50345), or as an array of probes on a solid
CC support.
XX
XX Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 19; Length 10;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 1 CTTCTCTTTT 10
RESULT 2
AAF33332
ID AAF33332 standard; DNA; 10 BP.
XX
XX AAF33332;
AC
XX
XX 23-MAR-2001 (first entry)
DT
XX
XX Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:71.
DE
XX
XX Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;
KW nor previously assigned open reading frame; nonannotated ORF; SAGE;
KW serial analysis of gene expression; antifungal; tag; identification;
XX linker; PCR primer; ds.
OS Saccharomyces cerevisiae.
XX
XX WO200077214-A2.
PN

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XX
XX 21-DEC-2000.
XX
XX 14-JUN-2000; 2000WO-US16223.
XX
XX 16-JUN-1999; 99US-0335032.
XX
XX (UYJO ) UNIV JOHNS HOPKINS.
XX
XX Velculescu V, Vogelstein B, Kinzler K;
XX
XX WPI; 2001-061874/07.
XX
XX Yeast gene coding sequences comprising NORF genes with serial analysis
PT of gene expression (SAGE) tags, useful for studying, monitoring and
PT affecting phases of the cell cycle -
XX
XX Claim 1; Page 23; 419pp; English.
XX
XX The present invention describes an isolated DNA molecule comprising a
CC coding sequence of a yeast gene selected from a group of 745 NORF (not
CC previously assigned open reading frame; or nonannotated ORF) genes
CC comprising a SAGE (serial analysis of gene expression) tag. Also
CC described are: (1) a method (M1) of using NORF genes to affect the cell
CC cycle comprising administering a NORF gene whose expression varies by at
CC least 10% between any two phases of the cell cycle selected from log
CC phase, S phase and G2/M; (2) a method (M2) for screening candidate
CC antifungal drugs comprising: (a) contacting a test substance with a
CC yeast cell; and (b) monitoring expression of a NORF gene whose
CC expression varies as in M1, where a test substance which modifies the
CC expression of the yeast gene is a candidate antifungal drug; (3) a method
CC (M3) for identifying human genes which are involved in cell cycle
CC progression comprising contacting human DNA with a probe which comprises
CC at least 10 contiguous nucleotides of a NORF gene whose expression varies
CC as in M1; and (4) a method (M4) for identifying a candidate drug as a
CC member of a class of drugs having a characteristic effect on gene
CC expression in a yeast cell comprising contacting a yeast cell with a
CC candidate drug and monitoring expression in the yeast cell of at least 1
CC NORF gene whose expression is affected by the class of drugs. The NORF
CC genes may be used to study, monitor and affect phases of the cell cycle,
CC the differentially expressed genes may be used as markers of phases of
CC the cell cycle. The methods may be used to identify candidate drugs which
CC affect the cell cycle and for identification of antifungal drugs.
CC AAF33268 to AAF4064 represent SAGE tags used in the exemplification of
CC the present invention. AAF33262 to AAF33267 represent linkers and PCR
CC primers used in the SAGE method, in the exemplification of the present
CC invention.
XX
XX Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 22; Length 10;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 1 CTTCTCTTTT 10
RESULT 3
AAF33921
ID AAF33921 standard; DNA; 10 BP.
XX
XX AAF33921;
AC
XX
XX 23-MAR-2001 (first entry)
DT
XX
XX Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:660.
DE
XX
XX Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;
KW nor previously assigned open reading frame; nonannotated ORF; SAGE;
KW serial analysis of gene expression; antifungal; tag; identification;
XX linker; PCR primer; ds.
KW

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XX OS Saccharomyces cerevisiae.  
 XX PN WO200077214-A2.  
 XX PD 21-DEC-2000.  
 XX PF 14-JUN-2000; 2000WO-US16223.  
 XX PR 16-JUN-1999; 99US-0335032.  
 XX PA (UYJO ) UNIV JOHNS HOPKINS.  
 XX PI Velculescu V, Vogelstein B, Kinzler K;  
 XX DR WPI; 2001-061874/07.  
 XX PT Yeast gene coding sequences comprising NORF genes with serial analysis  
 PT of gene expression (SAGE) tags, useful for studying, monitoring and  
 PT affecting phases of the cell cycle -  
 XX  
 PS Claim 1; Page 398; 419pp; English.  
 XX  
 CC The present invention describes an isolated DNA molecule comprising a  
 CC coding sequence of a yeast gene selected from a group of 745 NORF (not  
 CC previously assigned open reading frame; or nonannotated ORF) genes  
 CC comprising a SAGE (serial analysis of gene expression) tag. Also  
 CC described are: (1) a method (M1) of using NORF genes to affect the cell  
 CC cycle comprising administering a NORF gene whose expression varies by at  
 CC least 10% between any two phases of the cell cycle selected from log  
 CC phase, S phase and G2/M; (2) a method (M2) for screening candidate  
 CC antifungal drugs comprising: (a) contacting a test substance with a  
 CC yeast cell; and (b) monitoring expression of a NORF gene whose  
 CC expression varies as in M1, where a test substance which modifies the  
 CC expression of the yeast gene is a candidate antifungal drug; (3) a method  
 CC (M3) for identifying human genes which are involved in cell cycle  
 CC progression comprising contacting human DNA with a probe which comprises  
 CC at least 10 contiguous nucleotides of a NORF gene whose expression varies  
 CC as in M1; and (4) a method (M4) for identifying a candidate drug as a  
 CC member of a class of drugs having a characteristic effect on gene  
 CC expression in a yeast cell comprising contacting a yeast cell with a  
 CC candidate drug and monitoring expression in the yeast cell of at least 1  
 CC NORF gene whose expression is affected by the class of drugs. The NORF  
 CC genes may be used to study, monitor and affect phases of the cell cycle,  
 CC the differentially expressed genes may be used as markers of phases of  
 CC the cell cycle. The methods may be used to identify candidate drugs which  
 CC affect the cell cycle and for identification of antifungal drugs.  
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of  
 CC the present invention. AAF33262 to AAF33267 represent linkers and PCR  
 CC primers used in the SAGE method, in the exemplification of the present  
 CC invention.  
 XX  
 SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;  
 Query Match 100.0%; Score 10; DB 22; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 1 CTTCTCTTTT 10  
 RESULT 4  
 AAF33924  
 ID AAF33924 standard; DNA; 10 BP.  
 XX AAF33924;  
 XX  
 DT 23-MAR-2001 (first entry)  
 XX  
 DE Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:663.  
 XX

KW Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;  
 KW nor previously assigned open reading frame; nonannotated ORF; SAGE;  
 KW serial analysis of gene expression; antifungal; tag; identification;  
 KW linker; PCR primer; ds.  
 XX  
 OS Saccharomyces cerevisiae.  
 XX PN WO200077214-A2.  
 XX PD 21-DEC-2000.  
 XX PF 14-JUN-2000; 2000WO-US16223.  
 XX PR 16-JUN-1999; 99US-0335032.  
 XX PA (UYJO ) UNIV JOHNS HOPKINS.  
 XX PI Velculescu V, Vogelstein B, Kinzler K;  
 XX DR WPI; 2001-061874/07.  
 XX PT Yeast gene coding sequences comprising NORF genes with serial analysis  
 PT of gene expression (SAGE) tags, useful for studying, monitoring and  
 PT affecting phases of the cell cycle -  
 XX  
 PS Claim 1; Page 398; 419pp; English.  
 XX  
 CC The present invention describes an isolated DNA molecule comprising a  
 CC coding sequence of a yeast gene selected from a group of 745 NORF (not  
 CC previously assigned open reading frame; or nonannotated ORF) genes  
 CC comprising a SAGE (serial analysis of gene expression) tag. Also  
 CC described are: (1) a method (M1) of using NORF genes to affect the cell  
 CC cycle comprising administering a NORF gene whose expression varies by at  
 CC least 10% between any two phases of the cell cycle selected from log  
 CC phase, S phase and G2/M; (2) a method (M2) for screening candidate  
 CC antifungal drugs comprising: (a) contacting a test substance with a  
 CC yeast cell; and (b) monitoring expression of a NORF gene whose  
 CC expression varies as in M1, where a test substance which modifies the  
 CC expression of the yeast gene is a candidate antifungal drug; (3) a method  
 CC (M3) for identifying human genes which are involved in cell cycle  
 CC progression comprising contacting human DNA with a probe which comprises  
 CC at least 10 contiguous nucleotides of a NORF gene whose expression varies  
 CC as in M1; and (4) a method (M4) for identifying a candidate drug as a  
 CC member of a class of drugs having a characteristic effect on gene  
 CC expression in a yeast cell comprising contacting a yeast cell with a  
 CC candidate drug and monitoring expression in the yeast cell of at least 1  
 CC NORF gene whose expression is affected by the class of drugs. The NORF  
 CC genes may be used to study, monitor and affect phases of the cell cycle,  
 CC the differentially expressed genes may be used as markers of phases of  
 CC the cell cycle. The methods may be used to identify candidate drugs which  
 CC affect the cell cycle and for identification of antifungal drugs.  
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of  
 CC the present invention. AAF33262 to AAF33267 represent linkers and PCR  
 CC primers used in the SAGE method, in the exemplification of the present  
 CC invention.  
 XX  
 SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;  
 Query Match 100.0%; Score 10; DB 22; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 1 CTTCTCTTTT 10  
 RESULT 5  
 AAF34233  
 ID AAF34233 standard; DNA; 10 BP.  
 XX AAF34233;  
 XX

DT 23-MAR-2001 (first entry)  
 XX Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:972.  
 DE  
 XX  
 XX Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;  
 KW nor previously assigned open reading frame; nonannotated ORF; SAGE;  
 KW serial analysis of gene expression; antifungal; tag; identification;  
 KW linker; PCR primer; ds.  
 XX  
 XX Saccharomyces cerevisiae.  
 OS  
 XX WO200077214-A2.  
 PN  
 XX 21-DEC-2000.  
 XX  
 XX 14-JUN-2000; 2000WO-US16223.  
 PF  
 XX 16-JUN-1999; 99US-0335032.  
 PR  
 XX (UVOJO ) UNIV JOHNS HOPKINS.  
 PA  
 XX Velculescu V, Vogelstein B, Kinzler K;  
 PI WPI; 2001-061874/07.  
 XX  
 XX Yeast gene coding sequences comprising NORF genes with serial analysis  
 of gene expression (SAGE) tags, useful for studying, monitoring and  
 affecting phases of the cell cycle -  
 PT  
 XX Example; Page 34; 419pp; English.  
 XX  
 XX The present invention describes an isolated DNA molecule comprising a  
 coding sequence of a yeast gene selected from a group of 745 NORF (not  
 previously assigned open reading frame; or nonannotated ORF) genes  
 comprising a SAGE (serial analysis of gene expression) tag. Also  
 described are: (1) a method (M1) of using NORF genes to affect the cell  
 cycle comprising administering a NORF gene whose expression varies by at  
 least 10% between any two phases of the cell cycle selected from log  
 phase, S phase and G2/M; (2) a method (M2) for screening candidate  
 antifungal drugs comprising: (a) contacting a test substance with a  
 yeast cell; and (b) monitoring expression of a NORF gene whose  
 expression varies as in M1, where a test substance which modifies the  
 expression of the yeast gene is a candidate antifungal drug; (3) a method  
 (M3) for identifying human genes which are involved in cell cycle  
 progression comprising contacting human DNA with a probe which comprises  
 at least 10 contiguous nucleotides of a NORF gene whose expression varies  
 as in M1; and (4) a method (M4) for identifying a candidate drug as a  
 member of a class of drugs having a characteristic effect on gene  
 expression in a yeast cell comprising contacting a yeast cell with a  
 candidate drug and monitoring expression in the yeast cell of at least 1  
 NORF gene whose expression is affected by the class of drugs. The NORF  
 genes may be used to study, monitor and affect phases of the cell cycle,  
 the differentially expressed genes may be used as markers of phases of  
 the cell cycle. The methods may be used to identify candidate drugs which  
 affect the cell cycle and for identification of antifungal drugs.  
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of  
 the present invention. AAF33262 to AAF33267 represent linkers and PCR  
 CC primers used in the SAGE method, in the exemplification of the present  
 CC invention.  
 XX  
 XX Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;  
 SQ  
 Query Match 100.0%; Score 10; DB 22; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 DE |||||||  
 Db 1 CTTCTCTTTT 10  
 XX  
 RESULT 6  
 AAX14794/c

DT 23-MAR-2001 (first entry)  
 XX Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:972.  
 DE  
 XX  
 XX Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;  
 KW nor previously assigned open reading frame; nonannotated ORF; SAGE;  
 KW serial analysis of gene expression; antifungal; tag; identification;  
 KW linker; PCR primer; ds.  
 XX  
 XX Saccharomyces cerevisiae.  
 OS  
 XX WO200077214-A2.  
 PN  
 XX 21-DEC-2000.  
 XX  
 XX 14-JUN-2000; 2000WO-US16223.  
 PF  
 XX 16-JUN-1999; 99US-0335032.  
 PR  
 XX (UVOJO ) UNIV JOHNS HOPKINS.  
 PA  
 XX Velculescu V, Vogelstein B, Kinzler K;  
 PI WPI; 2001-061874/07.  
 XX  
 XX Yeast gene coding sequences comprising NORF genes with serial analysis  
 of gene expression (SAGE) tags, useful for studying, monitoring and  
 affecting phases of the cell cycle -  
 PT  
 XX Example; Page 34; 419pp; English.  
 XX  
 XX The present invention describes an isolated DNA molecule comprising a  
 coding sequence of a yeast gene selected from a group of 745 NORF (not  
 previously assigned open reading frame; or nonannotated ORF) genes  
 comprising a SAGE (serial analysis of gene expression) tag. Also  
 described are: (1) a method (M1) of using NORF genes to affect the cell  
 cycle comprising administering a NORF gene whose expression varies by at  
 least 10% between any two phases of the cell cycle selected from log  
 phase, S phase and G2/M; (2) a method (M2) for screening candidate  
 antifungal drugs comprising: (a) contacting a test substance with a  
 yeast cell; and (b) monitoring expression of a NORF gene whose  
 expression varies as in M1, where a test substance which modifies the  
 expression of the yeast gene is a candidate antifungal drug; (3) a method  
 (M3) for identifying human genes which are involved in cell cycle  
 progression comprising contacting human DNA with a probe which comprises  
 at least 10 contiguous nucleotides of a NORF gene whose expression varies  
 as in M1; and (4) a method (M4) for identifying a candidate drug as a  
 member of a class of drugs having a characteristic effect on gene  
 expression in a yeast cell comprising contacting a yeast cell with a  
 candidate drug and monitoring expression in the yeast cell of at least 1  
 NORF gene whose expression is affected by the class of drugs. The NORF  
 genes may be used to study, monitor and affect phases of the cell cycle,  
 the differentially expressed genes may be used as markers of phases of  
 the cell cycle. The methods may be used to identify candidate drugs which  
 affect the cell cycle and for identification of antifungal drugs.  
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of  
 the present invention. AAF33262 to AAF33267 represent linkers and PCR  
 CC primers used in the SAGE method, in the exemplification of the present  
 CC invention.  
 XX  
 XX Sequence 12 BP; 8 A; 0 C; 4 G; 0 U; 0 other;  
 SQ  
 Query Match 100.0%; Score 10; DB 20; Length 12;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 DE |||||||  
 Db 11 CTTCTCTTTT 2  
 XX  
 RESULT 7  
 AAX14859  
 ID AAX14859 standard; DNA; 12 BP.  
 XX  
 AC AAX14859;  
 XX  
 XX 24-MAR-1999 (first entry)  
 DT  
 XX Triple helix third strand of 23S rRNA gene nucleotides 214-225.  
 DE  
 XX Triple helix formation; DNA detection; triple helix; identification;  
 KW bacteria; oncogene; virus; ss.  
 XX  
 OS Synthetic.

ID AAX14794 standard; DNA; 12 BP.  
 XX  
 AC AAX14794;  
 XX  
 DT 24-MAR-1999 (first entry)  
 XX  
 DE Triple helix forming nucleotides 807-818 of Hepatitis B virus.  
 XX  
 KW Triple-helix forming region; Triplex formation; DNA detection;  
 KW identification; bacteria; oncogene; virus; ds.  
 XX  
 OS Hepatitis B virus.  
 XX  
 XX US5861244-A.  
 PN  
 XX 19-JAN-1999.  
 PD  
 XX 22-DEC-1993; 93US-0173489.  
 PF  
 XX 22-DEC-1993; 93US-0173489.  
 PR  
 XX 29-OCT-1992; 92US-0968436.  
 PR  
 XX (PROF-) PROFILE DIAGNOSTIC SCI INC.  
 PA  
 XX Hepburn AG, Wang C;  
 PI WPI; 1999-130384/11.  
 XX  
 XX Assay of genetic sequences based on triplex formation from double  
 stranded analyte - and hybrid of anchor and reporter sequences, with  
 reporter released if triplex formation occurs, used e.g. to identify  
 bacteria  
 PT  
 XX Disclosure; Columns 19-20; 168pp; English.  
 PS  
 XX The present sequence represents a potential triple-helix forming region.  
 CC It can be used to demonstrate the assay of the invention. The assay  
 CC comprises adding a sample containing double-stranded DNA test sequences,  
 CC e.g. containing the present sequence, to an aqueous medium containing at  
 CC least one complex of anchor DNA, attached to a solid support, and  
 CC reporter DNA, where either a part of the anchor DNA or reporter DNA is  
 CC designed to form a triple-strand structure with part of the test  
 CC sequence. Triplex formation results in displacement of the reporter DNA  
 CC which is detected as an indication of the presence of the DNA test  
 CC sequence. The method is used to detect DNA sequences, particularly for  
 CC identification of bacteria (by detecting genes for ribosomal RNA) in  
 CC clinical samples, but also detection of oncogenes and Hepatitis B virus.  
 XX  
 SQ Sequence 12 BP; 8 A; 0 C; 4 G; 0 U; 0 other;  
 Query Match 100.0%; Score 10; DB 20; Length 12;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 DE |||||||  
 Db 11 CTTCTCTTTT 2  
 XX  
 RESULT 7  
 AAX14859  
 ID AAX14859 standard; DNA; 12 BP.  
 XX  
 AC AAX14859;  
 XX  
 XX 24-MAR-1999 (first entry)  
 DT  
 XX Triple helix third strand of 23S rRNA gene nucleotides 214-225.  
 DE  
 XX Triple helix formation; DNA detection; triple helix; identification;  
 KW bacteria; oncogene; virus; ss.  
 XX  
 OS Synthetic.

```

OS Micrococcus luteus.
PN US5861244-A.
XX
PD 19-JAN-1999.
XX
PF 22-DEC-1993; 93US-0173489.
XX
PR 22-DEC-1993; 93US-0173489.
PR 29-OCT-1992; 92US-0969436.
XX
PA (PROF-) PROFILE DIAGNOSTIC SCI INC.
XX
PI Hepburn AG, Wang C;
XX
DR WPI; 1999-130384/11.
XX
PT Assay of genetic sequences based on triplex formation from double
PT stranded analyte - and hybrid of anchor and reporter sequences, with
PT reporter released if triplex formation occurs, used e.g. to identify
PT bacteria
XX
PS Disclosure; Columns 21-22; 168pp; English.
XX
CC The present sequence represents a polynucleotide that is able to
CC form a triple helix with a double stranded sequence. Cytosine bases
CC in the present can be replaced with 5-methylcytosine for increased
CC triplex stability. The present sequence is used in the assay of the
CC invention, where it can be part of the anchor DNA or reporter DNA
CC sequence. The assay comprises adding a sample containing double-stranded
CC DNA test sequences to an aqueous medium containing at least one complex
CC of anchor DNA, attached to a solid support, and reporter DNA, where
CC either a part of the anchor DNA or reporter DNA is designed to form
CC a triple-strand structure with part of the test sequence. Triplex
CC formation results in displacement of the reporter DNA which is
CC detected as an indication of the presence of the DNA test sequence.
CC The method is used to detect DNA sequences, particularly for
CC identification of bacteria (by detecting genes for ribosomal RNA) in
CC clinical samples, but also detection of oncogenes and Hepatitis B virus.
XX
SQ Sequence 12 BP; 0 A; 4 C; 0 G; 8 T; 0 other;
Query Match 100.0%; Score 10; DB 20; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 3 CTTCTCTTTT 12
RESULT 8
ABH67817
ID ABH67817 standard; DNA; 12 BP.
XX
AC ABH67817;
XX
DT 22-FEB-2002 (first entry)
XX
Oligonucleotide primer SEQ ID NO 267794 for detecting SNP TSC0000531.
DE
DE SNP: single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB00713.
XX
Set of oligonucleotides, useful for diagnosis and cell typing, is
designed to detect single nucleotide polymorphisms and cytosine
methylation status -
PT
PT methylation status -
PT
PT Claim 1; SEQ ID 380025; 29pp + Sequence Listing; German.
XX

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PR 07-APR-2000; 2000DE-1019173.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;
XX
DR WPI; 2001-657177/75.
XX
PT Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single nucleotide polymorphisms and cytosine
PT methylation status -
XX
PS Claim 1; SEQ ID 267794; 29pp + Sequence Listing; German.
XX
CC This invention describes novel oligonucleotide primers or peptide nucleic
CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
CC and cytosine methylation status in chemically pretreated genomic DNA. The
CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
CC range of diseases including immune system, gastrointestinal, respiratory,
CC central nervous system, cardiovascular and metabolic disorders. The
CC oligomers are also used for detecting cell type differentiation.
CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
CC ABT00010-ABT99989 represent the oligomers described in the invention.
CC NOTE: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 12 BP; 1 A; 4 C; 0 G; 7 T; 0 other;
Query Match 100.0%; Score 10; DB 23; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11
RESULT 9
ABI80052
ID ABI80052 standard; DNA; 12 BP.
XX
AC ABI80052;
XX
DT 22-FEB-2002 (first entry)
XX
Oligonucleotide primer SEQ ID NO 380025 for detecting SNP TSC0063600.
DE
DE SNP: single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB00713.
XX
PT 07-APR-2000; 2000DE-1019173.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;
XX
DR WPI; 2001-657177/75.
XX
PT Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single nucleotide polymorphisms and cytosine
PT methylation status -
XX
PS Claim 1; SEQ ID 380025; 29pp + Sequence Listing; German.
XX

```



XX This invention describes novel oligonucleotide primers or peptide nucleic  
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)  
 CC and cytosine methylation status in chemically pretreated genomic DNA. The  
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a  
 CC range of diseases including immune system, gastrointestinal, respiratory,  
 CC central nervous system, cardiovascular and metabolic disorders. The  
 CC oligomers are also used for detecting cell type differentiation.  
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and  
 CC AB100010-AB182073 represent the oligomers described in the invention.  
 CC NOTE: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 12 BP; 1 A; 4 C; 0 G; 7 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 12;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 3 CTTCTCTTTT 12

RESULT 10  
 ABC07094/c  
 ID ABC07094 standard; DNA; 13 BP.  
 AC ABC07094;  
 XX  
 DT 20-FEB-2002 (first entry)  
 XX  
 DE Oligonucleotide SEQ ID NO 7085 for detecting SNP TSC00002097.  
 XX  
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;  
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;  
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200177384-A2.  
 XX  
 PD 18-OCT-2001.  
 XX  
 PF 06-APR-2001; 2001WO-IB00713.  
 XX  
 PR 07-APR-2000; 2000DE-1019173.  
 XX  
 PA (EP1G-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K;  
 XX  
 DR WPI; 2001-657177/75.  
 XX  
 PT Set of oligonucleotides, useful for diagnosis and cell typing, is  
 PT designed to detect single nucleotide polymorphisms and cytosine  
 PT methylation status -  
 XX  
 PS Claim 1; SEQ ID 7085; 29pp + Sequence Listing; German.  
 CC This invention describes novel oligonucleotide primers or peptide nucleic  
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)  
 CC and cytosine methylation status in chemically pretreated genomic DNA. The  
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a  
 CC range of diseases including immune system, gastrointestinal, respiratory,  
 CC central nervous system, cardiovascular and metabolic disorders. The  
 CC oligomers are also used for detecting cell type differentiation.  
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and  
 CC AB100010-AB182073 represent the oligomers described in the invention.  
 CC NOTE: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 13 BP; 8 A; 0 C; 4 G; 1 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 13 CTTCTCTTTT 4

RESULT 11  
 ABC07095  
 ID ABC07095 standard; DNA; 13 BP.  
 XX  
 AC ABC07095;  
 XX  
 DT 20-FEB-2002 (first entry)  
 XX  
 DE Oligonucleotide SEQ ID NO 7086 for detecting SNP TSC00002097.  
 XX  
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;  
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;  
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200177384-A2.  
 XX  
 PD 18-OCT-2001.  
 XX  
 PF 06-APR-2001; 2001WO-IB00713.  
 XX  
 PR 07-APR-2000; 2000DE-1019173.  
 XX  
 PA (EP1G-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K;  
 XX  
 DR WPI; 2001-657177/75.  
 XX  
 PT Set of oligonucleotides, useful for diagnosis and cell typing, is  
 PT designed to detect single nucleotide polymorphisms and cytosine  
 PT methylation status -  
 XX  
 PS Claim 1; SEQ ID 7086; 29pp + Sequence Listing; German.  
 CC This invention describes novel oligonucleotide primers or peptide nucleic  
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)  
 CC and cytosine methylation status in chemically pretreated genomic DNA. The  
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a  
 CC range of diseases including immune system, gastrointestinal, respiratory,  
 CC central nervous system, cardiovascular and metabolic disorders. The  
 CC oligomers are also used for detecting cell type differentiation.  
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and  
 CC AB100010-AB182073 represent the oligomers described in the invention.  
 CC NOTE: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 13 BP; 1 A; 4 C; 0 G; 8 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 1 CTTCTCTTTT 10

RESULT 12

```

ABC53772/c
ID ABC53772 standard; DNA; 13 BP.
XX
AC ABC53772;
XX
DT 21-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 53789 for detecting SNP TSC0014813.
XX
KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB00713.
XX
PR 07-APR-2000; 2000DE-1019173.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;
XX
WPI; 2001-657177/75.
XX
Set of oligonucleotides, useful for diagnosis and cell typing, is
designed to detect single nucleotide polymorphisms and cytosine
methylation status -
XX
Claim 1; SEQ ID 53790; 29pp + Sequence Listing; German.
XX
This invention describes novel oligonucleotide primers or peptide nucleic
acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
and cytosine methylation status in chemically pretreated genomic DNA. The
oligonucleotides are used for diagnosis and/or prognosis of cancer and a
range of diseases including immune system, gastrointestinal, respiratory,
central nervous system, cardiovascular and metabolic disorders. The
oligonucleotides are also used for detecting cell type differentiation.
XX
ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
ABT00010-ABT99989 represent the oligomers described in the invention.
XX
NOTE: The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format from WIPO at
ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 13 BP; 10 A; 0 C; 3 G; 0 U; 0 other;
XX
Query Match 100.0%; Score 10; DB 23; Length 13;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1
XX
RESULT 13
ABC53773
ID ABC53773 standard; DNA; 13 BP.
XX
AC ABC53773;
XX
DT 21-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 53790 for detecting SNP TSC0014813.
XX
KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB00713.
XX
PR 07-APR-2000; 2000DE-1019173.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;
XX
WPI; 2001-657177/75.
XX
Set of oligonucleotides, useful for diagnosis and cell typing, is
designed to detect single nucleotide polymorphisms and cytosine
methylation status -
XX
Claim 1; SEQ ID 53789; 29pp + Sequence Listing; German.
XX
This invention describes novel oligonucleotide primers or peptide nucleic
acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
and cytosine methylation status in chemically pretreated genomic DNA. The
oligonucleotides are used for diagnosis and/or prognosis of cancer and a
range of diseases including immune system, gastrointestinal, respiratory,
central nervous system, cardiovascular and metabolic disorders. The
oligonucleotides are also used for detecting cell type differentiation.
XX
ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
ABT00010-ABT99989 represent the oligomers described in the invention.
XX
NOTE: The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format from WIPO at
ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 13 BP; 10 A; 0 C; 3 G; 0 U; 0 other;
XX
Query Match 100.0%; Score 10; DB 23; Length 13;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1
XX
RESULT 14
ABH25906/c
ID ABH25906 standard; DNA; 13 BP.
XX
AC ABH25906;
XX
DT 22-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 225883 for detecting SNP TSC0055065.
XX
KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB00713.
XX
PR 07-APR-2000; 2000DE-1019173.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;

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XX WP1; 2001-657177/75.  
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is  
 PT designed to detect single nucleotide polymorphisms and cytosine  
 PT methylation status -  
 XX PS Claim 1; SEQ ID 225883; 29pp + Sequence Listing; German.  
 XX This invention describes novel oligonucleotide primers or peptide nucleic  
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)  
 CC and cytosine methylation status in chemically pretreated genomic DNA. The  
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a  
 CC range of diseases including immune system, gastrointestinal, respiratory,  
 CC central nervous system, cardiovascular and metabolic disorders. The  
 CC oligomers are also used for detecting cell type differentiation.  
 CC NOTE: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.  
 XX SQ Sequence 13 BP; 8 A; 0 C; 3 G; 2 T; 0 other;  
 Query Match 100.0%; Score 10; DB 23; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 10 CTCTCTCTTT 1  
 RESULT 15  
 ID ABH25907 standard; DNA; 13 BP.  
 XX AC ABH25907;  
 XX DT 22-FEB-2002 (first entry)  
 XX DE Oligonucleotide SEQ ID NO 225884 for detecting SNP TSC0055065.  
 XX SNF: single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;  
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;  
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.  
 XX OS Homo sapiens.  
 XX WO200177384-A2.  
 XX PD 18-OCT-2001.  
 XX PF 06-APR-2001; 2001WO-IB00713.  
 XX PR 07-APR-2000; 2000DE-1019173.  
 XX PA (EP1G-) EPIGENOMICS AG.  
 XX PI Olek A, Piepenbrock C, Berlin K;  
 XX WP1; 2001-657177/75.  
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is  
 PT designed to detect single nucleotide polymorphisms and cytosine  
 PT methylation status -  
 XX PS Claim 1; SEQ ID 225884; 29pp + Sequence Listing; German.  
 XX This invention describes novel oligonucleotide primers or peptide nucleic  
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)  
 CC and cytosine methylation status in chemically pretreated genomic DNA. The  
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a

CC range of diseases including immune system, gastrointestinal, respiratory,  
 CC central nervous system, cardiovascular and metabolic disorders. The  
 CC oligomers are also used for detecting cell type differentiation.  
 CC AB000010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and  
 CC ABT00010-ABT2073 represent the oligomers described in the invention.  
 CC NOTE: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.  
 XX SQ Sequence 13 BP; 2 A; 3 C; 0 G; 8 T; 0 other;  
 Query Match 100.0%; Score 10; DB 23; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 4 CTCTCTCTTT 13  
 RESULT 16  
 ID ABV80474/c  
 XX ID ABV80474 standard; DNA; 17 BP.  
 XX AC ABV80474;  
 XX DT 03-JAN-2003 (first entry)  
 XX DE Human HTPL scanning oligonucleotide SEQ ID 1720.  
 XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
 XX OS Homo sapiens.  
 XX EP1229046-A2.  
 XX PD 07-AUG-2002.  
 XX PF 28-JAN-2002; 2002EP-0001167.  
 XX PR 30-JAN-2001; 2001WO-US000663.  
 XX PR 30-JAN-2001; 2001WO-US000664.  
 XX PR 30-JAN-2001; 2001WO-US000665.  
 XX PR 30-JAN-2001; 2001WO-US000667.  
 XX PR 30-JAN-2001; 2001WO-US000668.  
 XX PR 30-JAN-2001; 2001WO-US000669.  
 XX PR 23-MAY-2001; 2001US-0864761.  
 XX PR 09-OCT-2001; 2001US-0327898.  
 XX PA (AEON-) AEOMICA INC.  
 XX PI Zhan J;  
 XX WP1; 2002-676582/73.  
 XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX PS Example 2; Page 289; 718pp; English.  
 XX The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and AB898519 to AB898520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was

CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 0 C; 7 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 17 CTTCTCTTTT 8

RESULT 17

ABV80475/c  
 ID ABV80475 standard; DNA; 17 BP.

XX AC ABV80475;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1721.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 30-JAN-2001; 2001WO-US00669.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar

CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate, are  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 16 CTTCTCTTTT 7

RESULT 18

ABV80476/c

ID ABV80476 standard; DNA; 17 BP.

XX AC ABV80476;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1722.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001WO-US00669.

XX PR 09-OCT-2001; 2001US-0864761.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL

CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

SQ Sequence 17 BP; 9 A; 0 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 Db 15 CTTCTCTTTT 6

# RESULT 19

ABV80477/c  
 ID ABV80477 standard; DNA; 17 BP.

XX AC ABV80477;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1723.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV8759 to ABV8762 and ABV8519 to ABV8520). HTPL  
 CC has two isoforms, with a few single base pair differences between the

CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

SQ Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 14 CTTCTCTTTT 5

# RESULT 20

ABV80478/c

ID ABV80478 standard; DNA; 17 BP.

XX AC ABV80478;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1724.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like

CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX  
 SQ Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 13 CTTCTCTTTT 4

RESULT 21  
 ABV80479/c  
 ID ABV80479 standard; DNA; 17 BP.  
 XX  
 AC ABV80479;  
 XX  
 DT 03-JAN-2003 (first entry)  
 XX  
 DE Human HTPL scanning oligonucleotide SEQ ID 1725.  
 XX  
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP1229046-A2.  
 XX  
 PD 07-AUG-2002.  
 XX  
 PF 28-JAN-2002; 2002EP-0001167.  
 XX  
 PR 30-JAN-2001; 2001WO-US00663.  
 PR 30-JAN-2001; 2001WO-US00664.  
 PR 30-JAN-2001; 2001WO-US00665.  
 PR 30-JAN-2001; 2001WO-US00667.  
 PR 30-JAN-2001; 2001WO-US00668.  
 PR 30-JAN-2001; 2001WO-US00669.  
 PR 23-MAY-2001; 2001US-0864761.  
 PR 09-OCT-2001; 2001US-0327898.  
 XX  
 PA (AEOM-) AEOMICA INC.  
 XX  
 PI Zhan J;  
 XX  
 DR WPI; 2002-676582/73.  
 XX  
 PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX  
 PS Example 2; Page 290; 718pp; English.

XX  
 CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX  
 SQ Sequence 17 BP; 8 A; 1 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 12 CTTCTCTTTT 3

RESULT 22  
 ABV80480/c  
 ID ABV80480 standard; DNA; 17 BP.  
 XX  
 AC ABV80480;  
 XX  
 DT 03-JAN-2003 (first entry)  
 XX  
 DE Human HTPL scanning oligonucleotide SEQ ID 1726.  
 XX  
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP1229046-A2.  
 XX  
 PD 07-AUG-2002.  
 XX  
 PF 28-JAN-2002; 2002EP-0001167.  
 XX  
 PR 30-JAN-2001; 2001WO-US00663.  
 PR 30-JAN-2001; 2001WO-US00664.  
 PR 30-JAN-2001; 2001WO-US00665.  
 PR 30-JAN-2001; 2001WO-US00667.  
 PR 30-JAN-2001; 2001WO-US00668.  
 PR 30-JAN-2001; 2001WO-US00669.  
 PR 23-MAY-2001; 2001US-0864761.  
 PR 09-OCT-2001; 2001US-0327898.  
 XX  
 PA (AEOM-) AEOMICA INC.  
 XX  
 PI Zhan J;  
 XX  
 DR WPI; 2002-676582/73.  
 XX  
 PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 290; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like

CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL

CC has two isoforms, with a few single base pair differences between the

CC two. One of the single base pair changes introduces a premature stop

CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL

CC shares an overall structure organisation with the Patched protein. The

CC shared structural features strongly imply that HTPL plays a role similar

CC to that of Patched, and is a potential tumour suppressor. HTPL is

CC important in regulating male germ cell development, and the HTPL gene was

CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are

CC useful for diagnosing a disorder caused by mutation in HTPL, and in

CC therapy and manufacture of a medicament for treatment or prevention of

CC such disorder associated with decreased expression or activity of human

CC HTPL. Such disorders include disorders of testis, or adrenal, adult and

CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,

CC skeletal muscle or colon function. HTPL proteins and nucleic acids are

CC clinically useful diagnostic markers and potential therapeutic agents for

CC male infertility and cancer. The present oligonucleotide was used in an

CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 1 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 11 CTTCTCTTTT 2

RESULT 23

ABV80481/C

ID ABV80481 standard; DNA; 17 BP.

XX AC ABV80481;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1727.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;

XX KW human testis expressed Patched like protein; testis; adrenal; liver;

XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;

XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 30-JAN-2001; 2001WO-US00669.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AECOMICA INC.

XX PI Zhan J;

XX PS WPI; 2002-676582/73.

XX DR Novel isolated human testis expressed Patched like protein (HTPL),

XX PT

PT useful for identifying agonist and antagonist and specific binding

PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 290; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like

CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL

CC has two isoforms, with a few single base pair differences between the

CC two. One of the single base pair changes introduces a premature stop

CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL

CC shares an overall structure organisation with the Patched protein. The

CC shared structural features strongly imply that HTPL plays a role similar

CC to that of Patched, and is a potential tumour suppressor. HTPL is

CC important in regulating male germ cell development, and the HTPL gene was

CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are

CC useful for diagnosing a disorder caused by mutation in HTPL, and in

CC therapy and manufacture of a medicament for treatment or prevention of

CC such disorder associated with decreased expression or activity of human

CC HTPL. Such disorders include disorders of testis, or adrenal, adult and

CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,

CC skeletal muscle or colon function. HTPL proteins and nucleic acids are

CC clinically useful diagnostic markers and potential therapeutic agents for

CC male infertility and cancer. The present oligonucleotide was used in an

CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 1 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 10 CTTCTCTTTT 1

RESULT 24

ABV74761/C

ID ABS74761 standard; DNA; 17 BP.

XX AC ABS74761;

XX DT 24-DEC-2002 (first entry)

XX DE Human PAPP-Ea associated 17-mer SEQ ID 287.

XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;

XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;

XX KW dysgenetic pregnancy; primer; ss.

XX OS Homo sapiens.

XX PN US2002102252-A1.

XX PD 01-AUG-2002.

XX PF 06-APR-2001; 2001US-0827998.

XX PR 26-MAY-2000; 2000US-207456P.

XX PA (GUYV/) GU Y.

XX PA (SHAN/) SHANNON M E.

XX PI Gu Y, Shannon ME;

XX DR WPI; 2002-697817/75.

XX PT New isolated nucleic acid encoding an isoform of human pregnancy

XX PT associated plasma protein E, for preventing or aborting pregnancy -

XX PS Example 2; Page 113; 353pp; English.

XX CC This invention describes a novel isolated nucleic acid that encodes

CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 10 A; 2 C; 4 G; 1 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 17 CTTCTCTTTT 8

RESULT 25  
 ABS74762/c  
 ID ABS74762 standard; DNA; 17 BP.

AC ABS74762;  
 XX  
 XX  
 DT 24-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 17-mer SEQ ID 288.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.

XX Homo sapiens.

XX US2002102252-A1.

XX 01-AUG-2002.

XX 06-APR-2001; 2001US-0827998.

XX 26-MAY-2000; 2000US-207456P.

XX (GUYY/) GU Y.

XX (SHAN/) SHANNON M E.

XX Gu Y, Shannon ME;

XX WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy -  
 XX Example 2; Page 113; 353pp; English.

CC This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 10 A; 2 C; 4 G; 1 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 16 CTTCTCTTTT 7

RESULT 26  
 ABS74763/c  
 ID ABS74763 standard; DNA; 17 BP.

AC ABS74763;  
 XX  
 XX  
 DT 24-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 17-mer SEQ ID 289.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.

XX Homo sapiens.

XX US2002102252-A1.

XX 01-AUG-2002.

XX 06-APR-2001; 2001US-0827998.

XX 26-MAY-2000; 2000US-207456P.

XX (GUYY/) GU Y.

XX (SHAN/) SHANNON M E.

XX Gu Y, Shannon ME;

XX WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 XX Example 2; Page 113; 353pp; English.

CC This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 9 A; 2 C; 4 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 15 CTTCTCTTTT 6



```

RESULT 27
ABS74764/c
ID ABS74764 standard; DNA; 17 BP.
XX
XX AC ABS74764;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 290.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX
XX PA (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX DR WPI; 2002-697817/75.
XX
XX PT New isolated nucleic acid encoding an isoform of human pregnancy
XX PT associated plasma protein E, for preventing or aborting pregnancy -
XX
XX PS Example 2; Page 113; 353pp; English.
XX
XX CC This invention describes a novel isolated nucleic acid that encodes
XX CC one of three new isoforms of human pregnancy associated plasma protein E,
XX CC hPAPP-E. The products of the invention have abortive and contraceptive
XX CC activity and can be used for gene therapy or in a vaccine. The nucleic
XX CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX CC used in pharmaceutical compositions or vaccines for preventing or
XX CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX CC antibodies can be used to assess the expression levels of PAPP-E isoform
XX CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX CC antenatally. This sequence represents an oligomer used in scanning the
XX CC human PAPP-E genes described in the disclosure of the invention.
XX
XX SQ Sequence 17 BP; 10 A; 1 C; 4 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
DB 14 CTCTCTCTTT 5

RESULT 28
ABS74765/c
ID ABS74765 standard; DNA; 17 BP.
XX
XX AC ABS74765;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 291.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW

```

```

KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX
XX PA (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX DR WPI; 2002-697817/75.
XX
XX PT New isolated nucleic acid encoding an isoform of human pregnancy
XX PT associated plasma protein E, for preventing or aborting pregnancy -
XX
XX PS Example 2; Page 113; 353pp; English.
XX
XX CC This invention describes a novel isolated nucleic acid that encodes
XX CC one of three new isoforms of human pregnancy associated plasma protein E,
XX CC hPAPP-E. The products of the invention have abortive and contraceptive
XX CC activity and can be used for gene therapy or in a vaccine. The nucleic
XX CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX CC used in pharmaceutical compositions or vaccines for preventing or
XX CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX CC antibodies can be used to assess the expression levels of PAPP-E isoform
XX CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX CC antenatally. This sequence represents an oligomer used in scanning the
XX CC human PAPP-E genes described in the disclosure of the invention.
XX
XX SQ Sequence 17 BP; 11 A; 1 C; 4 G; 1 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
DB 13 CTCTCTCTTT 4

RESULT 29
ABS74766/c
ID ABS74766 standard; DNA; 17 BP.
XX
XX AC ABS74766;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 292.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX

```

PA (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX  
 PI Gu Y, Shannon ME;  
 XX  
 DR WPI; 2002-697817/75.  
 XX  
 PT New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 XX  
 XX Example 2; Page 113; 353pp; English.  
 XX  
 CC This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 17 BP; 10 A; 1 C; 4 G; 2 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 12 CTCTCTCTTT 3  
 RESULT 30  
 ABS74767/C  
 ID ABS74767 standard; DNA; 17 BP.  
 XX  
 AC ABS74767;  
 XX  
 DT 24-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 17-mer SEQ ID 293.  
 XX  
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN US2002102252-A1.  
 XX  
 PD 01-AUG-2002.  
 XX  
 PF 06-APR-2001; 2001US-0827998.  
 XX  
 PR 26-MAY-2000; 2000US-207456P.  
 XX  
 PA (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX  
 PI Gu Y, Shannon ME;  
 XX  
 DR WPI; 2002-697817/75.  
 XX  
 PT New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 XX  
 XX Example 2; Page 113; 353pp; English.  
 XX  
 CC This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 17 BP; 10 A; 1 C; 4 G; 2 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 12 CTCTCTCTTT 3  
 RESULT 31  
 ABS74768/C  
 ID ABS74768 standard; DNA; 17 BP.  
 XX  
 AC ABS74768;  
 XX  
 DT 24-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 17-mer SEQ ID 294.  
 XX  
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN US2002102252-A1.  
 XX  
 PD 01-AUG-2002.  
 XX  
 PF 06-APR-2001; 2001US-0827998.  
 XX  
 PR 26-MAY-2000; 2000US-207456P.  
 XX  
 PA (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX  
 PI Gu Y, Shannon ME;  
 XX  
 DR WPI; 2002-697817/75.  
 XX  
 PT New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 XX  
 XX Example 2; Page 113; 353pp; English.  
 XX  
 CC This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 17 BP; 10 A; 0 C; 4 G; 3 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 11 CTCTCTCTTT 2

CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 17 BP; 9 A; 1 C; 4 G; 3 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 10 CTTCTCTTTT 1  
 |||||  
 RESULT 32  
 ABX94543  
 ID ABX94543 standard; DNA; 19 BP.  
 XX  
 AC ABX94543;  
 XX  
 DT 13-JUN-2003 (first entry)  
 XX  
 DE 23S/16S rRNA detecting probe SEQ ID 12.  
 XX  
 KW Detection; probe; contaminant; drinking water; Legionella; coliform;  
 KW faecal streptococci; soil; sputum; biopsy; urine; food; pharmaceutical;  
 KW cosmetic; fluorescent in situ hybridisation; FISH; ss.  
 XX  
 OS Streptococcus sp.  
 XX  
 PN WO2002102824-A2.  
 XX  
 PD 27-DEC-2002.  
 XX  
 PF 19-JUN-2002; 2002WO-EP06809.  
 XX  
 PR 19-JUN-2001; 2001DE-1029411.  
 PR 11-DEC-2001; 2001DE-1060666.  
 XX  
 PA (VERM-) VERMICON AG.  
 PI Beinfuhr C, Snaldr J;  
 XX  
 DR WPI; 2003-167479/16.  
 XX  
 PT New oligonucleotides, useful for detecting bacteria that may  
 PT contaminate drinking water, provide quick results for many species in  
 PT parallel -  
 XX  
 PS Claim 8; Page 12; 53pp; German.  
 XX  
 CC This invention describes novel oligonucleotide probes used to detect  
 CC contaminant bacteria that may be present in drinking water. The probes  
 CC can detect bacteria (especially Legionella, faecal streptococci and  
 CC coliforms) that may contaminate drinking water in environmental samples  
 CC (water or soil), clinical samples (sputum, biopsies, urine etc.), in  
 CC bathing and drinking water and in foods, pharmaceuticals and cosmetics,  
 CC by in situ hybridisation. The probes combine the advantages of  
 CC fluorescent in situ hybridisation with those of culture methods. Only a  
 CC relatively short culture step is required; analysis takes 24-48 hours  
 CC (contrast many days for conventional methods) and all relevant bacteria  
 CC can be tested simultaneously. The oligonucleotides can differentiate  
 CC between species of the same genus and are easy to use, allowing simple  
 CC analysis of a large number of samples. ABX94532-ABX94578 represent the  
 CC oligonucleotide probes described in the invention.  
 XX  
 SQ Sequence 19 BP; 1 A; 7 C; 2 G; 9 T; 0 other;  
 Query Match 100.0%; Score 10; DB 25; Length 19;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 1 CTTCTCTTTT 10  
 |||||

Db 4 CTTCTCTTTT 13  
 RESULT 33  
 AAT76779  
 ID AAT76779 standard; DNA; 20 BP.  
 XX  
 AC AAT76779;  
 XX  
 DT 15-SEP-1997 (first entry)  
 XX  
 DE Staphylococcus aureus exfoliative toxin A gene PCR primer ETA-B.  
 XX  
 KW Asymmetric polymerase chain reaction; nucleic acid amplification;  
 KW PCR; detection; assay; exfoliative toxin A; ETA; skin lesion;  
 KW competitive primer; capture probe; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN US5627054-A.  
 XX  
 PD 06-MAY-1997.  
 XX  
 PF 05-APR-1996; 96US-0628417.  
 XX  
 PR 05-APR-1996; 96US-0628417.  
 XX  
 PA (USSA ) US SEC OF ARMY.  
 XX  
 PI Gillespie D;  
 XX  
 DR WPI; 1997-271311/24.  
 XX  
 PT Quantitative nucleic acid amplification - by competitor primer  
 PT asymmetric polymerase chain reaction  
 XX  
 PS Example 1; Column 5; 9pp; English.  
 XX  
 CC In a specific example of a novel process for amplifying an amount  
 CC (known or unknown) of a double-stranded nucleic acid segment to produce  
 CC single-stranded nucleic acid in an amount that is proportional to the  
 CC starting amount of the nucleic acid, the Staphylococcus aureus  
 CC exfoliative toxin A (ETA) gene was used as the DNA template. The  
 CC region comprising nucleotides 165-436 was amplified by symmetric,  
 CC asymmetric or competitor primer asymmetric PCR using the primers  
 CC ETA-A2 and ETA-B (see AAT76778 and AAT76779). For asymmetric PCR, the  
 CC amount of primer ETA-B was reduced and for competitor primer  
 CC asymmetric PCR a competitor primer ETA-B2 (see AAT76780) was added  
 CC with upstream primer ETA-A2 after the initial cycling reaction. PCR  
 CC products containing ETA-specific sequences were detected  
 CC radioactively by a capture system which employed a bifunctional  
 CC capture probe ETA-CP (see AAT76781 and AAT76782). ETA-CP was designed  
 CC to capture the amplified sense strand onto capture membranes  
 CC through hybridisation between the first 40 nucleotides of ETA-CP  
 CC and nucleotides 321-360 of the ETA gene and through hybridisation  
 CC of the poly(dA) tail on ETA-CP with poly(dT) tails on the capture  
 CC membranes. A radioactively labelled "label probe" (see AAT76783),  
 CC complementary to nucleotides 389-410 of the ETA gene was used to  
 CC detect the amplicons. Results showed that hybridisation of the  
 CC capture probe and label probe to the denatured symmetric PCR  
 CC product was much less efficient than hybridisation to the  
 CC single-stranded PCR products of the asymmetric and competitor  
 CC primer asymmetric reactions.  
 XX  
 SQ Sequence 20 BP; 4 A; 5 C; 3 G; 8 T; 0 other;  
 Query Match 100.0%; Score 10; DB 18; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 8 CTTCTCTTTT 17  
 |||||

RESULT 34  
AAZ04007  
ID AAZ04007 standard; DNA; 20 BP.  
XX AC  
XX AAZ04007;  
XX  
DT 07-OCT-1999 (first entry)  
XX  
DE PCR primer used to amplify an ORF of Chlamydia trachomatis.  
XX  
KW Vaccine; eye disease; conventional trachoma; nonendemic trachoma;  
KW paratrachoma; inclusion conjunctivitis; genital disease; perihhepatitis;  
KW nongonococcal urethritis; epididymitis; cervicitis; salpingitis; PCR primer;  
KW Bartholinitis; pneumopathy; venereal lymphogranulomatosis; ss.  
XX  
OS Synthetic.  
OS Chlamydia trachomatis.  
XX  
PN WO9928475-A2.  
XX  
PD 10-JUN-1999.  
XX  
XX 27-NOV-1998; 98WO-IB01939.  
XX  
PR 04-NOV-1998; 98US-0107077.  
PR 28-NOV-1997; 97FR-0015041.  
PR 17-DEC-1997; 97FR-0016034.  
XX  
XX (GEST ) GENSET.  
XX  
PA Griffais R;  
PI  
XX WPI; 1999-371125/31.  
XX  
XX Genome sequence of Chlamydia trachomatis  
PT  
XX Disclosure; Page 1653; 1755pp; English.  
XX  
XX PCR primers AAZ01426-206209 were used to amplify open reading frames  
CC (ORFs) of the genome of Chlamydia trachomatis (see AAZ01425). These ORFs  
CC encode polypeptides (see AAX36754-Y37949) which can be used as vaccines  
CC against Chlamydia trachomatis. Antisense and ribozyme sequences  
CC can also be used to control growth of the microorganism. Chlamydia  
CC trachomatis is responsible for a large number of diseases, e.g. eye  
CC diseases such as conventional trachoma, nonendemic trachoma,  
CC paratrachoma, and inclusion conjunctivitis; genital diseases such as  
CC nongonococcal urethritis, epididymitis, cervicitis, salpingitis,  
CC perihhepatitis, Bartholinitis, pneumopathy in breast feeding infants,  
CC and venereal lymphogranulomatosis. The polypeptides of the  
CC invention may be of use in treating these diseases.  
XX  
SQ Sequence 20 BP; 0 A; 7 C; 3 G; 10 T; 0 other;  
Query Match 100.0%; Score 10; DB 20; Length 20;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
Db 3 CTTCTCTTTT 12  
RESULT 35  
AAX97056/c  
ID AAX97056 standard; DNA; 20 BP.  
XX AC  
XX AAX97056;  
XX  
DT 13-SEP-1999 (first entry)  
XX  
DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;  
KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;  
KW vaccine; neutralising epitope; PCR primer; ss.  
XX  
OS Synthetic.  
OS Chlamydia pneumoniae.  
XX  
PN WO9927105-A2.  
XX  
XX 03-JUN-1999.  
XX  
PD 20-NOV-1998; 98WO-IB01890.  
XX  
PF 04-NOV-1998; 98US-0107078.  
PR 21-NOV-1997; 97FR-0014673.  
XX  
XX (GEST ) GENSET.  
XX  
PA Griffais R;  
PI  
XX WPI; 1999-357842/30.  
XX  
XX Genome sequence of Chlamydia pneumoniae  
PT  
XX Page 1874; Disclosure; 1912pp; English.  
XX  
XX AAX91991-X97517 represent PCR primers used to amplify open reading  
CC frames and other nucleic acid sequences from the genome of  
CC Chlamydia pneumoniae (see AAX91990). C. pneumoniae causes respiratory  
CC disease such as pneumonia and bronchitis and is thought to be a  
CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent  
CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded  
CC by the open reading frames of the C. pneumoniae genome (see AAX94584-  
CC AAX35879) can be used in immunogenic compositions as vaccines. Vectors  
CC containing C. pneumoniae nucleotide sequences can also be used as  
CC immunogenic compositions, especially where the vector directs the  
CC expression of a neutralising epitope of C. pneumoniae.  
XX  
SQ Sequence 20 BP; 8 A; 4 C; 4 G; 4 T; 0 other;  
Query Match 100.0%; Score 10; DB 20; Length 20;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
Db 17 CTTCTCTTTT 8  
RESULT 36  
AAL51617/c  
ID AAL51617 standard; DNA; 20 BP.  
XX AC  
XX AAL51617;  
XX  
DT 17-APR-2003 (first entry)  
XX  
XX Human interferon alpha 2 sequencing primer #5.  
DE  
XX Human; ss; cellular proliferation inhibitor; interferon alpha 2;  
KW single nucleotide polymorphism; SNP; cancer; tumour; metabolic disease;  
KW cardiovascular disease; infectious disease; immunological disease; HIV;  
KW central nervous system disease; wound healing; chemotherapy side effect;  
KW anaemia; osteoporosis; gastrointestinal disease; venereal disease; AIDS;  
KW obesity; hepatitis; infectious pneumonia; Alzheimer's disease; allergy;  
KW Parkinson's disease; multiple sclerosis; schizophrenia; depression;  
KW graft versus host disease; asthma; psoriasis; rheumatoid arthritis;  
KW Crohn's disease; ulcerative colitis; genital wart; sequencing; primer.  
XX  
OS Homo sapiens.  
XX  
PN EP1236800-A2.

XX PD 04-SEP-2002.

XX PF 01-MAR-2002; 2002EP-0290515.

XX PR 01-MAR-2001; 2001FR-0002843.

XX PA (GENO-) GENODYSSEE.

XX PI Escary J;

XX PI WPI; 2003-185789/19.

XX DR

XX PT An isolated polynucleotide encoding interferon alpha 2 containing

XX PT single nucleotide polymorphisms is useful in treating disease -

XX PS Example 4; Page 21; 42pp; English.

XX CC The invention comprises the amino acid and coding sequence of the human

XX CC interferon alpha 2 protein. The invention further relates to the

XX CC identification of single nucleotide polymorphisms (SNPs) within the human

XX CC interferon alpha 2 gene. The DNA and protein sequences of the invention

XX CC are useful for the treatment of: cancer; tumours; cardiovascular

XX CC diseases; metabolic diseases; infectious diseases; central nervous system

XX CC diseases; immunological diseases; wound healing; chemotherapy side

XX CC effects; anaemia; osteoporosis; Gastrointestinal diseases; venereal

XX CC diseases; obesity; hepatitis; HIV/AIDS; infectious pneumonias;

XX CC Alzheimer's disease; Parkinson's disease; multiple sclerosis;

XX CC schizophrenia; depression; graft versus host disease; allergies; asthma;

XX CC psoriasis; rheumatoid arthritis; Crohn's disease; ulcerative colitis; and

XX CC genital warts. The present DNA sequence represents a primer that was used

XX CC to sequence the human interferon alpha 2 gene.

XX SQ

Query Match 100.0%; Score 10; DB 25; Length 20;

Best Local Similarity 100.0%; Pred No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 20 CTTCTCTTTT 11

RESULT 37

AAx60140/c

ID AAX60140 standard; DNA; 21 BP.

XX AC AAX60140;

XX DT 05-AUG-1999 (first entry)

XX DE PCR primer used to amplify Mycoplasma hyopneumoniae P102 protein DNA.

XX KW P102 protein; vaccine; antigen; diagnosis; swine; immunisation;

XX KW enzootic pneumonia; PCR primer; ss.

XX OS Synthetic.

XX PN WO926664-A1.

XX PD 03-JUN-1999.

XX PF 24-NOV-1998; 98WO-US25044.

XX PR 26-NOV-1997; 97US-0066565.

XX PA (IOWA) UNIV IOWA STATE RES FOUND INC.

XX PI Hsu T, Minion PC;

XX DR WPI; 1999-357741/30.

PT Recombinant antigenic Mycoplasma hyopneumoniae protein

XX PS Example 2; Page 23; 45pp; English.

XX CC PCR primers AAX60140-41 were used to amplify DNA encoding a Mycoplasma

XX CC hyopneumoniae P102 protein clone. The P102 protein and its fragments

XX CC are used in vaccines to protect against enzootic pneumonia,

XX CC particularly in swine. Recombinant P102 polypeptides may be used as

XX CC antigens for diagnostic purposes to determine whether or not a

XX CC biological test sample contains M. hyopneumoniae antigens or

XX CC antibodies. The P102 polypeptides or DNA sequences may also be used

XX CC for immunising or protecting non-human animals, preferably swine,

XX CC against M. hyopneumoniae infections, particularly enzootic pneumonia.

XX SQ Sequence 21 BP; 12 A; 0 C; 7 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 21;

Best Local Similarity 100.0%; Pred No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 15 CTTCTCTTTT 6

RESULT 38

AAV08126/c

ID AAV08126 standard; DNA; 21 BP.

XX AC AAV08126;

XX DT 22-JAN-1999 (first entry)

XX DE Primer Vbeta14 for T cell receptor V region.

XX KW PCR primer; T-cell receptor; TCR; V region; immune response; arthritis;

XX KW somatic homologous recombination; hypervariable region;

XX KW spectratype determination; autoimmune response; multiple sclerosis;

XX KW myasthenia gravis; muscular dystrophy; graft-infiltrating lymphocyte;

XX KW tumour-infiltrating lymphocyte; ss.

XX OS Synthetic.

XX OS Mammalia.

XX PN US5837447-A.

XX PD 17-NOV-1998.

XX PF 19-APR-1994; 94US-0229528.

XX PR 19-APR-1994; 94US-0229528.

XX PR 15-APR-1992; 92US-0868569.

XX XX (BLOO-) BLOOD CENT RES FOUND INC.

XX PA Gorski J;

XX PI WPI; 1999-023435/02.

XX DR

XX PT Monitoring immune responses by analysing amplified B or T-cell

XX PT nucleic acid - using primers specific for variable and constant or

XX PT junction region gene segments, with separation of products by

XX PT length, especially to monitor auto:immunity

XX PS Claim 22; Column 38; 26pp; English.

XX CC This sequence represents a primer for the T cell receptor (TCR) variable

XX CC region and is used in the method of the invention. The method is for

XX CC monitoring an immune response that involves somatic homologous

XX CC recombination between elements of at least two segments associated with a

XX CC hypervariable region, and comprises: (a) providing a polynucleotide

XX CC sample from B- or T-cells, and amplifying it with: (i) a primer specific

XX CC for a variable gene segment; and (ii) a primer specific for a constant or

CC joining gene segment to produce amplification products (AP) that can be  
 CC resolved at a difference in size of 2 or 3 bp; (c) separating the AP  
 CC according to length; (d) detecting the range of lengths in the separated  
 CC products to produce a 'spectratype' of the subject's immune response; and  
 CC (e) comparing the spectratype with a predetermined standard to determine  
 CC immune status or to monitor immune response. The method is specifically  
 CC used to monitor autoimmune responses (including relapses), i.e. to  
 CC identify the predominant TCR in sites of autoimmune activity (e.g. in  
 CC arthritis, multiple sclerosis, myasthenia gravis and muscular dystrophy)  
 CC or present in graft-infiltrating (in cases of organ rejection) or  
 CC tumour-infiltrating lymphocytes. As each gene rearrangement is unique,  
 CC each complementarity determining region 3 is a specific molecular  
 CC fingerprint of the lymphocyte that generates it, and immune responses can  
 CC be correlated with an increase in a particular TCR or immunoglobulin.  
 CC Specific determination of two V beta families may be done simultaneously.  
 XX Sequence 21 BP; 8 A; 3 C; 7 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 18 CTTCTCTTTT 9

RESULT 39  
 AAA95613/c  
 ID AAA95613 standard; DNA; 21 BP.

AC AAA95613;

DT 31-JAN-2001 (first entry)  
 DE TCR Vbeta 14 subfamily probe VB14-1.

XX Detection; diagnostic; Kawasaki disease; T-cell; PCR primer; probe;  
 KW gene expression; ss.

XX Homo sapiens.

PN JP2000157297-A.

XX 13-JUN-2000.

PF 01-DEC-1998; 98JP-0341661.

PR 01-DEC-1998; 98JP-0341661.

XX (SHIO) SHIONOGI & CO LTD.

XX WPI; 2000-477722/42.

PT Detection of Kawasaki disease factor, useful for the diagnosis of  
 PT Kawasaki disease, comprises detecting an increase in Vbeta6.5 positive  
 PT T-cells -

PS Example 1; Page 9; 36pp; Japanese.

XX The invention relates to a method of detecting Kawasaki disease by  
 CC detecting an increase in Vbeta6.5 or Vbeta6.5/Vbeta2.1 positive T-cells.  
 CC The sequences AAA95531-A95626 represent primers and probes used to PCR  
 CC amplify and detect the level of expression of Valpha and Vbeta genes  
 CC in T-cells in Kawasaki disease.

SQ Sequence 21 BP; 8 A; 3 C; 7 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 18 CTTCTCTTTT 9  
 |||||

RESULT 40  
 AAH27157/c  
 ID AAH27157 standard; DNA; 21 BP.

XX AAH27157;

AC AAH27157;

DT 08-AUG-2001 (first entry)

DE Downstream PCR primer for amplification of PDAT gene.

XX Yeast; ARE 1; YCR048w; transgenic plant; oil production; acyl-CoA;  
 KW fatty acid production; triacylglycerol; oil crop; rape; sunflower; PDAT;  
 KW oil palm; soy; maize; oat; potato; sugar beet; turnip; PCR primer; ss.

XX Saccharomyces cerevisiae.

PN WO200134814-A1.

XX 17-MAY-2001.

PF 10-NOV-2000; 2000WO-SE02216.

XX 12-NOV-1999; 93EP-0850169.

PR 12-NOV-1999; 99US-0164859.

XX (SCAN-) SCANBI SCANDINAVIAN BIOTECHNOLOGY RES AB.

XX Banas A, Sandager L, Stahl U, Dahlqvist A, Lenman M, Ronne H;  
 PI Styrmne S;

XX WPI; 2001-329086/34.

XX Transforming oil-producing organisms with a gene encoding an  
 PT acyl-CoA:diacylglycerol acyltransferase, useful to generate  
 PT agricultural crops with higher triacylglycerol content -

PS Example 1; Page 6; 30pp; English.

XX This invention relates to the use of a novel enzyme in the production of  
 CC an oil-producing organism. The enzyme catalyses the transfer of a fatty  
 CC acid from acyl-CoA to diacylglycerol to produce triacylglycerol,  
 CC resulting in an increased oil content. Sequences AAH27155 and AAH97263  
 CC represent the Saccharomyces cerevisiae ARE1 coding and protein sequence  
 CC respectively. ARE1 is used in the transformation or the organism of the  
 CC invention. The invention is used to increase the oil content of oil crops  
 CC such as rape, sunflower and oil palm, and other crops such as soy, maize,  
 CC oat, potato, sugar beet and turnips. The invention could also be used to  
 CC produce triacylglycerols in microorganisms. The present sequence  
 CC represents a PCR primer used to amplify the Saccharomyces cerevisiae PDAT  
 CC gene. The primer and PCR product are used in an example illustrating that  
 CC triacylglycerol accumulation is reduced in yeast cells that lack the ARE1  
 CC gene. The primer is specifically used in the production of mutant yeast  
 CC strains.

SQ Sequence 21 BP; 13 A; 3 C; 5 G; 0 U; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||

Db 17 CTTCTCTTTT 8

RESULT 41  
 AA895067/c  
 ID AA895067 standard; DNA; 21 BP.

XX

```

AC AAS95067;
XX
DT 13-FEB-2002 (first entry)
XX
DE Human otoferlin exon PCR primer #32.
XX
KW Human; mouse; otoferlin; OTOF; brain; auditory function; PCR primer;
KW autosomal nonsyndromic prelingual deafness; DFNB9; ss.
XX
OS Homo sapiens.
XX
PN WO200170972-A2.
XX
PD 27-SEP-2001.
XX
PF 23-MAR-2001; 2001WO-IB00578.
XX
PR 24-MAR-2000; 2000US-191738P.
XX
PA (INSP ) INST PASTEUR.
PA (CNRS ) CNRS CENT NAT RECH SCI.
XX
PI Yasunaga S, Grati M, Cohen-Salmon M, El Amraoui A, Petit C;
PI Weil D;
XX
DR WPI; 2001-611499/70.
XX
XX Novel human gene Otoferlin, underlying an autosomal recessive
PT nonsyndromic prelingual deafness, DFNB9, and proteins encoded by the
PT gene, implicated in deafness -
XX
PS Claim 25; Page 17; 99pp; English.
XX
XX The invention relates to a purified polynucleotide (I) encoding a protein
CC sequence (II) encoded by a novel human gene, otoferlin (OTOF) or
CC the long human otoferlin isoform in brain. (I) was identified as
CC underlying an autosomal nonsyndromic prelingual deafness DFNB9, and is
CC thus useful for detecting deafness disease in humans and for
CC characterising the functions of proteins and genes encoding them in
CC auditory function. AAS95022-AAS95248 represent human and mouse
CC otoferlin coding sequences, PCR primers and related sequences of the
CC invention.
XX
SQ Sequence 21 BP; 10 A; 1 C; 8 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db |||||
10 CTTCTCTTTT 1

RESULT 42
AAX14678
ID AAX14678 standard; DNA; 22 BP.
XX
AC AAX14678;
XX
DT 24-MAR-1999 (first entry)
XX
DE Triple helix forming nucleotides 9-30 of gamma-crystallin gene.
XX
KW Triple-helix forming region; Triplex formation; DNA detection;
KW identification; bacteria; oncogene; virus; ds.
XX
OS Homo sapiens.
XX
PN US5861244-A.
XX
PD 19-JAN-1999.
XX

Query Match 100.0%; Score 10; DB 20; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db |||||
6 CTTCTCTTTT 15

RESULT 43
AAC66394/c
ID AAC66394 standard; DNA; 22 BP.
XX
AC AAC66394;
XX
DT 26-FEB-2001 (first entry)
XX
DE Human 3-hydroxyacyl-CoA-dehydratase cDNA specific PCR primer.
XX
KW Human; 3-hydroxyacyl-CoA-dehydratase; HCDase; hypothalamus;
KW PCR primer; ss.
XX
OS Homo sapiens.
XX
PN CN1263157-A.
XX
PD 16-AUG-2000.
XX
PF 17-FEB-2000; 2000CN-0111690.
XX
PR 17-FEB-2000; 2000CN-0111690.
XX
PA (NANF-) NANFANG RES CENT STATE HUMAN GENE GROUP.
XX
PI Li N, Qian B, Peng Y;
XX
DR WPI; 2000-639262/62.
XX
PT New human hydroxybutyryl coenzyme A dehydratase protein and its coding
PT sequence -

```

```

PF 22-DEC-1993; 93US-0173489.
XX
PR 22-DEC-1993; 93US-0173489.
PR 29-OCT-1992; 92US-0968436.
XX
PA (PROF-) PROFILE DIAGNOSTIC SCI INC.
XX
PI Hepburn AG, Wang C;
XX
DR WPI; 1999-130384/11.
XX
XX Assay of genetic sequences based on triplex formation from double
PT stranded analyte - and hybrid of anchor and reporter sequences, with
PT reporter released if triplex formation occurs, used e.g. to identify
PT bacteria
XX
PS Disclosure; Columns 15-16; 168pp; English.
XX
XX The present sequence represents a potential triple-helix forming region.
CC It can be used to demonstrate the assay of the invention. The assay
CC comprises adding a sample containing double-stranded DNA test sequences,
CC e.g. containing the present sequence, to an aqueous medium containing at
CC least one complex of anchor DNA, attached to a solid support, and
CC reporter DNA, where either a part of the anchor DNA or reporter DNA is
CC designed to form a triple-strand structure with part of the test
CC sequence. Triplex formation results in displacement of the reporter DNA
CC which is detected as an indication of the presence of the DNA test
CC sequence. The method is used to detect DNA sequences, particularly for
CC identification of bacteria (by detecting genes for ribosomal RNA) in
CC clinical samples, but also detection of oncogenes and Hepatitis B virus.
XX
SQ Sequence 22 BP; 1 A; 5 C; 0 G; 16 T; 0 other;

```

XX Example 1; Page 11; 21pp; Chinese.

PS This invention relates to a new human 3-hydroxyacyl-CoA-dehydratase

XX (HCDase). The protein is expressed in normal hypothalamic tissue in

CC humans. The invention includes human HCDase nucleotide and amino acid

CC sequences, a method for the preparation of the protein and a method for

CC detecting human HCDase nucleic acid and protein sequences in a sample.

CC The present sequence represents a PCR primer specific for human HCDase

CC cDNA.

XX

XX Sequence 22 BP; 10 A; 2 C; 8 G; 2 T; 0 other;

QQ

Query Match 100.0%; Score 10; DB 21; Length 22;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 16 CTTCTCTTTT 7

RESULT 44

ABL35686

ID ABL35686 standard; DNA; 22 BP.

XX

AC ABL35686;

XX

DT 04-APR-2002 (first entry)

XX

DE Immunostimulatory oligonucleotide SEQ ID NO: 612.

XX

KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;

KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;

KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;

KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;

KW antiinflammatory; antibacterial; ss.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT misc\_RNA 1..22

FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to

FT form RNA or DNA/RNA hybrids. Thymidine is linked to at

FT least one other base through a ribose sugar"

XX

PN WO200193902-A2.

XX

PD 13-DEC-2001.

XX

PF 07-JUN-2001; 2001WO-US18276.

XX

PR 07-JUN-2000; 2000US-209797P.

XX

PA (BIOS-) BIOSYNEXUS INC.

XX

PI Mond JJ, Flora M, Klinman DM;

XX

DR WPI; 2002-130570/17.

XX

PT New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing

PT cytokines, particularly for treating diseases, e.g. cancer, allergy or

PT HIV infection -

XX

PS Example 11; Page 63; 68pp; English.

XX

CC The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and

CC a DNA region. The composition is useful for enhancing an immune response

CC or inducing cytokines. It can be used as a vaccine adjuvant and in

CC treating diseases, including pathogenic infection, (non-)malignant

CC tumors (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC (e.g. allergic rhinitis, hay fever, or food allergies), Lyme disease,

CC hepatitis, HIV or malaria. The composition is also useful for treating,

CC preventing or ameliorating the symptoms resulting from exposure to a

CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence

CC is an immunostimulatory oligonucleotide described in the exemplification

XX of the invention.

XX

XX Sequence 22 BP; 1 A; 4 C; 3 G; 14 T; 0 other;

QQ

Query Match 100.0%; Score 10; DB 24; Length 22;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 7 CTTCTCTTTT 16

RESULT 45

ABL35687

ID ABL35687 standard; DNA; 22 BP.

XX

AC ABL35687;

XX

DT 04-APR-2002 (first entry)

XX

DE Immunostimulatory oligonucleotide SEQ ID NO: 613.

XX

KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;

KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;

KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;

KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;

KW antiinflammatory; antibacterial; ss.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT misc\_RNA 1..22

FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to

FT form RNA or DNA/RNA hybrids. Thymidine is linked to at

FT least one other base through a ribose sugar"

XX

PN WO200193902-A2.

XX

PD 13-DEC-2001.

XX

PF 07-JUN-2001; 2001WO-US18276.

XX

PR 07-JUN-2000; 2000US-209797P.

XX

PA (BIOS-) BIOSYNEXUS INC.

XX

PI Mond JJ, Flora M, Klinman DM;

XX

DR WPI; 2002-130570/17.

XX

PT New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing

PT cytokines, particularly for treating diseases, e.g. cancer, allergy or

PT HIV infection -

XX

PS Example 11; Page 63; 68pp; English.

XX

CC The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and

CC a DNA region. The composition is useful for enhancing an immune response

CC or inducing cytokines. It can be used as a vaccine adjuvant and in

CC treating diseases, including pathogenic infection, (non-)malignant

CC tumors (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies



CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,  
 CC hepatitis, HIV or malaria. The composition is also useful for treating,  
 CC preventing or ameliorating the symptoms resulting from exposure to a  
 CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence  
 CC is an immunostimulatory oligonucleotide described in the exemplification  
 CC of the invention.

XX Sequence 22 BP; 1 A; 4 C; 4 G; 13 T; 0 other;  
 SQ

Query Match 100.0%; Score 10; DB 24; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 Db 7 CTTCTCTTTT 16

RESULT 46  
 ABL35688  
 ID ABL35688 standard; DNA; 22 BP.  
 XX  
 AC ABL35688;  
 XX  
 DT 04-APR-2002 (first entry)  
 XX  
 DE Immunostimulatory oligonucleotide SEQ ID NO: 614.  
 XX  
 KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;  
 KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;  
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;  
 KW immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;  
 KW antiinflammatory; antibacterial; ss.  
 XX  
 OS Synthetic.  
 XX  
 FH Key Location/Qualifiers  
 FT misc\_RNA  
 FT 1..22  
 FT /tag= a  
 FT /note= "optionally thymidine is replaced by uracil to  
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at  
 FT least one other base through a ribose sugar"  
 XX  
 XX WO200193902-A2.  
 XX  
 PD 13-DEC-2001.  
 XX  
 PF 07-JUN-2001; 2001WO-US18276.  
 XX  
 PR 07-JUN-2000; 2000US-209797P.  
 XX  
 PA (BIOS-) BIOSYNEXUS INC.  
 XX  
 PI Mond JJ, Flora M, Klinman DM;  
 XX  
 WI; 2002-130570/17.  
 XX  
 DR New immunostimulatory compositions comprising RNA/DNA hybrid  
 XX oligonucleotides, useful for enhancing an immune response or inducing  
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or  
 PT HIV infection -  
 XX  
 PS Example 11; Page 63; 68pp; English.  
 XX  
 CC The present invention relates to an immunostimulatory composition, which  
 CC comprises at least one oligonucleotide comprising both an RNA region and  
 CC a DNA region. The composition is useful for enhancing an immune response  
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in  
 CC treating diseases, including pathogenic infection, (non-)malignant  
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or  
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies  
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,  
 CC hepatitis, HIV or malaria. The composition is also useful for treating,

CC preventing or ameliorating the symptoms resulting from exposure to a  
 CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence  
 CC is an immunostimulatory oligonucleotide described in the exemplification  
 CC of the invention.

XX Sequence 22 BP; 1 A; 4 C; 2 G; 15 T; 0 other;  
 SQ

Query Match 100.0%; Score 10; DB 24; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 Db 7 CTTCTCTTTT 16

RESULT 47  
 AAX10022/c  
 ID AAX10022 standard; DNA; 23 BP.  
 XX  
 AC AAX10022;  
 XX  
 DT 24-MAR-1999 (first entry)  
 XX  
 DE Human biallelic polymorphic marker downstream primer #328.  
 XX  
 KW Polymorphism; biallelic; human; forensic; paternity testing; disease;  
 KW detection; phenotypic typing; characteristic; infection; hereditary;  
 KW autoimmune disease; cancer; inflammation; drug; therapy; medicament;  
 XX treatment; marker; primer; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 FN WO9820165-A2.  
 XX  
 PD 14-MAY-1998.  
 XX  
 PF 05-NOV-1997; 97WO-US20313.  
 XX  
 PR 06-NOV-1996; 96US-0030455.  
 XX  
 PA (WHEED ) WHITEHEAD INST BIOMEDICAL RES.  
 XX  
 PI Hudson T, Lander ES, Wang D;  
 XX  
 WI; 1998-286974/25.  
 XX  
 DR New isolated nucleic acid segments from the human genome - used for  
 PT determining polymorphic forms for use in e.g. forensics, paternity  
 PT testing or phenotypic typing for disease  
 XX  
 PS Claim 16; Page 92; 310pp; English.  
 XX  
 CC AAX09121-X10268 are allele-specific oligonucleotide primers used in the  
 CC isolation of various biallelic polymorphic markers found in the human  
 CC genome (represented in AAX10269-X12937). These primers can be used in a  
 CC method for determining polymorphic forms in an individual for use in  
 CC e.g. forensics, paternity testing or for phenotypic typing for diseases  
 CC such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan syndrome,  
 CC muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease, familial  
 CC hypercholesterolemia, polycystic kidney disease, hereditary  
 CC spherocytosis, von Willebrand's disease, tubercular sclerosis, hereditary  
 CC haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos  
 CC syndrome, osteogenesis imperfecta, acute intermittent porphyria,  
 CC autoimmune diseases, inflammation, cancer, diseases of the nervous  
 CC system, infection by pathogenic microorganisms, and characteristics such  
 CC as longevity, appearance (e.g. baldness, obesity), strength, speed,  
 CC endurance, fertility, and susceptibility or receptivity to particular  
 CC drugs or therapeutic treatments. The isolated polymorphic nucleic acid  
 CC segments can also be used to produce medicaments for the treatment or  
 CC prophylaxis of such diseases.

```

SQ Sequence 23 BP; 16 A; 0 C; 7 G; 0 U; 0 other;
  Query Match      100.0%; Score 10; DB 19; Length 23;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 48
AAQ50920/c
ID AAQ50920 standard; DNA; 24 BP.
XX
AC AAQ50920;
XX
DT 25-MAR-2003 (updated)
DT 19-MAY-1994 (first entry)
XX
DE T-cell antigen receptor V-beta14 PCR primer.
XX
XX RT-PCR; polymerase chain reaction; amplification; SSCP;
KW single-strand conformation polymorphism; variable domain;
KW subtype beta 14; ss.
XX
OS Synthetic.
XX
PN WO9322455-A1.
XX
PD 11-NOV-1993.
XX
PF 30-APR-1993; 93WO-JP00577.
XX
PR 30-APR-1992; 92JP-0111467.
PR 31-JUL-1992; 92JP-0205054.
XX
PA (LTLI-) LTT INST CO LTD.
PA (TAIS) TAISHO PHARM CO LTD.
XX
PI Ikeda Y, Mizushima Y, Nishioka K, Sakoda H, Yamamoto K;
XX
XX WPI; 1993-368813/46.
XX
PT Detection of expression of T-cell antigen receptor gene - in
PT cancer, viral or immune disease patients, by polymerase chain
PT reaction amplification of the gene and SSCP analysis
XX
PS Claim 4; Page 13; 47pp; Japanese.
XX
CC Primers corresp. to DNA coding for part of the beta-chain of the T
CC cell antigen receptor (pref. the variable region primers AAQ50905-
CC AAQ50926) are used in PCR to amplify the T cell antigen receptor gene.
CC The amplified gene is detected by the single-strand conformation
CC polymorphism method using hybridisation probes corresp. to the
CC beta-chain J domain (see AAQ50928-Q50940).
CC (Updated on 25-MAR-2003 to correct FN field.)
XX
SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
  Query Match      100.0%; Score 10; DB 14; Length 24;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 49
AAAT10390/c
ID AAAT10390 standard; cDNA; 24 BP.
XX

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AC AAT10390;
XX
DT 02-APR-1996 (first entry)
XX
DE T-cell receptor primer Vbeta14.
XX
XX T cell receptor; beta chain; variable region; rheumatoid arthritis;
KW synovial joint fluid; PCR; amplification; primer; immunogen; vaccine;
KW immune disease; ss.
XX
OS Synthetic.
XX
PN WO9523164-A1.
XX
PD 31-AUG-1995.
XX
PF 23-FEB-1995; 95WO-EP00670.
XX
PR 23-FEB-1994; 94EP-0200454.
XX
PA (ALKU ) AKZO NOBEL NV.
XX
PI Graus JPM, Rijnders AWM, Van Der Maaden JM;
XX
XX WPI; 1995-311502/40.
XX
PT Peptide contained in the variable region of a T-cell receptor beta
PT chain - specifically associated with immune disease, esp. rheumatoid
PT arthritis
XX
PS Example 2; Page 24; 55pp; English.
XX
CC The primers AAT10352-97 were used to PCR amplify the T cell receptor
CC beta chain variable regions from T cell culture clones, isolated from
CC the synovial joint fluid of 11 patients suffering from rheumatoid
CC arthritis. The coding sequences were shown to contain the nucleotide
CC sequence AAT07409. The encoded polypeptide can be used as an immunogenic
CC cpd. for the detection of or predisposition to an immune disease, or for
CC use as a vaccine for prevention or treatment of an immune disease. This
CC primer amplifies the variable region from the 14.1 family of clones.
XX
SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
  Query Match      100.0%; Score 10; DB 16; Length 24;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 50
AAT98116/c
ID AAT98116 standard; DNA; 24 BP.
XX
AC AAT98116;
XX
DT 13-MAR-1998 (first entry)
XX
DE Primer V-beta(14) for T-cell receptor beta chain variable region.
XX
KW Antibody; T-cell receptor; beta chain; human immunodeficiency virus;
KW HIV; blood; attenuation; primer; PCR; amplification; variable region;
KW constant region; TCR; ss.
XX
OS Synthetic.
OS Homo sapiens.
XX
PN US5665355-A.
XX
PD 09-SEP-1997.
XX

```

PF 07-JUN-1995; 95US-0488212.  
 XX  
 PR 09-NOV-1992; 92US-0973485.  
 PR 18-OCT-1994; 94US-0408011.  
 PR 07-JUN-1995; 95US-0488212.  
 XX  
 PA (CONS-) CONSORZIO BIOTECNOLOGIE.  
 XX  
 XX Primi D;  
 XX WPI; 1997-456759/42.  
 XX  
 PT Removal of T-cell receptor-specific antibody from blood of  
 PT HIV-infected person - by extracorporeal blood treatment, to  
 PT attenuate or avert development of AIDS from HIV infection  
 XX  
 PS Example 1; Column 11; 43pp; English.  
 XX  
 CC The invention relates to a method for removing an antibody specific for  
 CC TCR-V beta (T-cell receptor V beta protein) from an HIV-infected person  
 CC by removing blood from the person, removing the antibody from the blood,  
 CC and reintroducing the blood into the person, thus allowing attenuation  
 CC or aversion of immunodeficiency. The primers AAT98100-T98150 are used  
 CC to check the efficiency of removal by detecting expression of the  
 CC TCR-V-beta and V-alpha genes in a blood sample after treatment. This  
 CC primer is targeted to the variable region sequence and can be used in  
 CC the amplification with primer AAT98100.  
 XX  
 SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 18; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 18 CTCTCTCTTT 9  
 RESULT 51  
 AAX85959/c  
 ID AAX85959 standard; DNA; 24 BP.  
 XX  
 AC AAX85959;  
 XX  
 DT 13-SEP-1999 (first entry)  
 XX  
 DE PCR primer used to amplify T cell receptor beta-chain cDNA.  
 XX  
 KW Acquired immune deficiency syndrome; free antibody; paratope; epitope;  
 KW T cell receptor variable beta region; TCR-V beta region; binding agent;  
 KW CD4+ T cell; HIV; PCR primer; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 PN US5928642-A.  
 XX  
 PD 27-JUL-1999.  
 XX  
 PF 18-OCT-1994; 94US-0408011.  
 XX  
 PR 09-NOV-1992; 92US-0973485.  
 PR 18-OCT-1994; 94US-0408011.  
 XX  
 PA (CONS-) CONSORZIO BIOTECNOLOGIE.  
 XX  
 XX Primi D;  
 XX WPI; 1999-429481/36.  
 DR  
 XX  
 PT Diagnosis and treatment of acquired immune deficiency syndrome  
 XX

PS Example 1; Column 11; 42pp; English.  
 XX  
 CC The specification describes a method for the diagnosis and treatment of  
 CC acquired immune deficiency syndrome, in a person having free antibodies  
 CC which have a paratope capable of binding to an epitope of a T cell  
 CC receptor variable beta (TCR-V beta) region. The method comprises  
 CC administering a binding agent homologous with the TCR-V beta  
 CC epitope. The binding agent is useful in assays for detecting various  
 CC CD4+ T cell subpopulations which carry particular V beta components.  
 CC The binding agent is also useful in the treatment of people infected  
 CC with HIV where it is able to remove an antibody able to bind with an  
 CC epitope on a TCR-V beta cell in the blood of an infected person. PCR  
 CC primers AAX85943-71 represent T cell receptor beta-chain primers used  
 CC in the course of the invention.  
 XX  
 SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 20; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 Db 18 CTCTCTCTTT 9  
 RESULT 52  
 AAX88135/c  
 ID AAX88135 standard; DNA; 24 BP.  
 XX  
 AC AAX88135;  
 XX  
 DT 09-SEP-1999 (first entry)  
 XX  
 DE T cell receptor beta chain primer V-beta14.  
 XX  
 KW T cell receptor; beta chain; primer; antibody; paratope; AIDS; vaccine;  
 KW epitope; TCR-V beta; immunogenic; anti-idiotypic; antiviral; detection;  
 KW CD4+ cell subpopulation; acquired immune deficiency syndrome; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN US5925513-A.  
 XX  
 PD 20-JUL-1999.  
 XX  
 PF 07-JUN-1995; 95US-0488209.  
 XX  
 PR 09-NOV-1992; 92US-0973485.  
 PR 18-OCT-1994; 94US-0408011.  
 PR 07-JUN-1995; 95US-0488209.  
 XX  
 PA (CONS-) CONSORZIO BIOTECNOLOGIE.  
 XX  
 XX Primi D;  
 XX WPI; 1999-418267/35.  
 DR  
 XX  
 PT Diagnosis and treatment of acquired immune deficiency syndrome onset  
 XX  
 PS Example 1; Column 11-12; 42pp; English.  
 XX  
 CC This invention describes novel method for binding free antibodies having  
 CC a paratope specific to an epitope on a T cell receptor (TCR-V beta)  
 CC while providing an immunogenic substance able to raise anti-idiotypic  
 CC antibodies which bind to free antibodies bound at the same paratope  
 CC specific to the epitope on the TCR-V beta and introducing this into a  
 CC person to raise anti-idiotypic antibodies. The products of the invention  
 CC have antiviral activity and can be used in vaccines. The specific  
 CC antibody binding affinities are useful in assays which detect the  
 CC presence of CD4+ cell subpopulations carrying particular V beta  
 CC components of the TCR-V beta in people infected with acquired immune  
 CC deficiency syndrome (AIDS). AAX88119-X88169 represents primers used in

CC the method of the invention.

SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 24;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 18 CTTCTCTTTT 9

RESULT 53

AAI99892/c

ID AAI99892 standard; DNA; 24 BP.

XX

AC AAI99892;

XX 30-JAN-2002 (first entry)

XX Human dihydroorotase 11 PCR primer SEQ ID NO 4.

XX Human; dihydroorotase 11; cytostatic; virucidal; immunomodulatory;

KW antiinflammatory; haemostatic; malignant tumour; HIV; infection;

KW human immunodeficiency virus; immunological disease; gene therapy;

KW PCR primer; ss.

XX Homo sapiens.

OS WO200173054-A1.

PN 04-OCT-2001.

XX 26-MAR-2001; 2001WO-CN00421.

XX 27-MAR-2000; 2000CN-0115157.

PR (SHAN-) SHANGHAI BIOWINDOW GENE DEV INC.

PI Mao Y, Xie Y;

XX WPI; 2001-602867/68.

DR New human dihydroorotase 11 for diagnosing and treating malignant

PT tumour, haemopathy, human immunodeficiency virus infection, immunological

PT diseases and various inflammations

XX Example 2; Page 17; 34pp; Chinese.

XX The invention relates to human dihydroorotase 11 with cytostatic,

CC virucidal, immunomodulatory, antiinflammatory and haemostatic

CC activity. The protein and encoding polynucleotide are used in diagnosis

CC and treatment of malignant tumour, haemopathy, human immunodeficiency

CC virus (HIV) infection, immunological diseases and various

CC inflammations. The polynucleotide is useful in gene therapy. The present

CC sequence is that of a PCR primer, useful to the invention.

XX

SQ Sequence 24 BP; 8 A; 3 C; 8 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 24;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 10 CTTCTCTTTT 1

RESULT 54

AAH19640/c

ID AAI99892 standard; DNA; 24 BP.

XX

AC

XX

DT

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KW

KW

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OS

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PF

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PR

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PI

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DR

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PT

PT

PT

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PS

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CC

KW multiple sclerosis; human; reverse transcriptase; RT-PCR; primer; ss.  
 XX Homo sapiens.  
 OS  
 XX US2002107388-A1.  
 PN  
 XX 08-AUG-2002.  
 PD  
 XX 10-MAY-2001; 2001US-0853830.  
 PF  
 XX 12-MAY-2000; 2000US-203984P.  
 PR  
 XX (VAND/) VANDENBARK A A.  
 PA  
 XX Vandenbark AA;  
 PI  
 XX WPI; 2002-697882/75.  
 DR  
 XX Identifying a T cell receptor variable gene expressed by target T cells  
 PT in an individual is useful to identify disease-associated T cells for  
 PT design of individualised therapies, particularly for autoimmune disease  
 PT -  
 XX  
 PS Example 2; Page 11; 20pp; English.  
 CC The invention relates to a method for identifying a T cell receptor  
 CC variable (TCRV) gene expressed by target T cells in an individual,  
 CC comprising determining expression of TCRV genes by activated T cells from  
 CC the individual and determining regulatory activity elicited in response  
 CC to TCRV peptides from the individual. A preferentially expressed TCRV  
 CC gene whose TCRV peptide elicits low T cell regulatory activity is  
 CC identified as a variable gene expressed by target T cells. The method is  
 CC used to identify disease-associated T cells in an individual so that  
 CC individualised therapies can be designed to prevent or treat the disease,  
 CC particularly an autoimmune disease, especially multiple sclerosis. This  
 CC sequence represents a reverse transcriptase PCR (RT-PCR) primer used in  
 CC analysis of expression of DNA encoding TCR variable beta (BV) peptides.  
 XX  
 SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 DB 18 CTTCTCTTTT 9  
 RESULT 56  
 ABK67667/C  
 ID ABK67667 standard; DNA; 24 BP.  
 XX  
 AC ABK67667;  
 AC  
 XX 02-JUL-2002 (first entry)  
 DT  
 DE B. subtilis knock-out PCR primer KO murA R1.  
 XX  
 KW Antibacterial; antiprotozoal; antifungal; cell growth inhibitor;  
 KW protozoa; bacterium; fungus; amoeba; mycoplasma; murA;  
 KW folA; ylbD; primer; ss.  
 XX  
 OS Bacillus subtilis.  
 OS  
 XX WO200216940-A2.  
 PN  
 XX 28-FEB-2002.  
 PD  
 XX 23-AUG-2001; 2001WO-US26322.  
 PF  
 XX 23-AUG-2000; 2000US-226896P.  
 PR  
 XX

(GENO-) GENOME THERAPEUTICS CORP.  
 PA Sulavik M, Ling LL, Opperman T, Moir DT, Bunker C;  
 PI  
 XX WPI; 2002-329705/36.  
 DR  
 XX Identification of a molecular target of a cell growth inhibitor,  
 PT comprises target prediction processes to identify at least one  
 PT modulated gene or gene product -  
 XX  
 PS Example 2; Page 97; 157pp; English.  
 CC The invention relates to a method of identifying a molecular target of a  
 CC cell growth inhibiting compound by: (a) identifying a compound or  
 CC composition inhibiting growth in a population of cells; (b) performing  
 CC target prediction processes to identify a modulated gene or gene product;  
 CC and (c) comparing by target prediction processes with the gene or gene  
 CC product identified by other prediction processes. The method is used for  
 CC identifying the molecular target of a cell growth inhibitor. The cells  
 CC are selected from organisms such as protozoa, bacterium, fungus, amoeba  
 CC and mycoplasma. The method reaps the benefits of both cell-based and  
 CC target-based screening. The method relies first on identification of  
 CC compounds with good whole cell activity and then provides methods to  
 CC identify the target of the compounds. Each of the methods is successful  
 CC in isolation, but in combination of at least two (preferably at least  
 CC three) provides a higher success in identifying the molecular target.  
 CC ABK67622-ABK67679 represent PCR primers used in examples which  
 CC demonstrate the method of the invention.  
 XX  
 SQ Sequence 24 BP; 9 A; 4 C; 6 G; 5 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 DB 16 CTTCTCTTTT 7  
 RESULT 57  
 ABA04339/C  
 ID ABA04339 standard; DNA; 24 BP.  
 XX  
 AC ABA04339;  
 AC  
 XX 05-MAR-2002 (first entry)  
 DT  
 DE Human BOLA structural domain zinc finger protein 37 PCR primer 1.  
 XX  
 KW Human; BOLA structural domain zinc finger protein 37; malignant tumour;  
 KW nosohaemia; HIV infection; immunological disease; inflammation;  
 KW PCR primer; ss.  
 XX  
 OS Homo sapiens.  
 OS  
 XX CN1307053-A.  
 PN  
 XX 08-AUG-2001.  
 PD  
 XX 28-JAN-2000; 2000CN-0111611.  
 PF  
 XX 28-JAN-2000; 2000CN-0111611.  
 PR  
 XX (BODA-) BODAO GENE TECH CO LTD SHANGHAI.  
 PA  
 XX Mao Y, Xie Y;  
 PI  
 XX WPI; 2002-062745/09.  
 DR  
 XX Polypeptide-human BOLA structural domain zinc finger protein 37 and  
 PT polynucleotide for said polypeptide -  
 XX

PS Example 3; Page 17 (Disclosure); 35pp; Chinese.  
XX  
CC The present invention describes human BOLA structural domain zinc finger  
CC protein 37. (i). (I) can be used in the treatment of various diseases,  
CC such as malignant tumour, nosohaemia, HIV infection, immunological  
CC diseases and inflammations. The present sequence represents a PCR primer  
CC for human BOLA structural domain zinc finger protein 37, which is used  
CC in an example from the present invention.  
XX  
SQ Sequence 24 BP; 12 A; 4 C; 4 G; 4 T; 0 other;  
XX  
Query Match 100.0%; Score 10; DB 24; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
QY 1 CTTCTCTTTT 10  
DB 18 CTTCTCTTTT 9  
XX  
RESULT 59  
ABL45589/c  
ID ABL45589 standard; DNA; 24 BP.  
XX  
AC ABL45589;  
XX  
DT 11-APR-2002 (first entry)  
XX  
DE Human chromosome 21q22.1 PCR primer SEQ ID NO:2633.  
XX  
KW Human; chromosome 1p36-35; chromosome 21q22.1; genetic analysis;  
KW genome; PCR primer; ss.  
XX  
OS Homo sapiens.  
XX  
PN JP2001321190-A.  
XX  
PD 20-NOV-2001.  
XX  
PF 12-MAR-2001; 2001JP-0068285.  
XX  
PR 10-MAR-2000; 2000JP-0066716.  
XX  
PA (RIKA) RIKAGAKU KENKYUSHO.  
PA (GENO-) GENOTEX YG.  
XX  
DR WPI; 2002-144136/19.  
XX  
PT Arraying genome clones -  
XX  
PS Claim 6; Page 57; 528pp; Japanese.  
XX  
CC The present invention describes a method of arraying genome clones. The  
CC method comprises: (a) clones of the genomic libraries contained in  
CC multiwell plates numbered for discrimination are mixed in each of the  
CC multiwell plates; (b) a primer designed based on the chromosome marker  
CC sequence is added to the mixture to carry out an amplification reaction;  
CC (c) a signal corresponding to the marker is detected from the resultant  
CC amplified product to specify the discrimination Nos. of the multiwell  
CC plates containing the clones having said marker sequence; (d) the order  
CC of the markers is changed so that the same discrimination Nos. succeed to  
CC the maximum in the specified discrimination Nos. to array the multiwell  
CC plates; (e) the clones in the multiwell plates of the specified  
CC discrimination Nos. are mixed respectively in each wells of longitudinal  
CC and lateral directions; (f) the mixed clones are cultured and the  
CC resultant cultures are amplified by using the above primer; (g) signals  
CC are detected from the amplified products; (h) the clones in the multiwell  
CC plates are specified from the detected result; and (i) the clones are  
CC reconstituted as the positions on the chromosome and arrayed. The  
CC microarray is useful for gene analysis. ABL42957 to ABL45322 represent  
CC PCR primers for human chromosome 1p36-35 DNA, and ABL45323 to ABL45634  
CC represent PCR primers for human chromosome 21q22.1, which are  
CC specifically claimed for use in the present invention.  
XX

XX  
SQ Sequence 24 BP; 12 A; 2 C; 6 G; 4 T; 0 other;  
XX  
Query Match 100.0%; Score 10; DB 24; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
QY 1 CTTCTCTTTT 10  
DB 19 CTTCTCTTTT 10  
XX  
RESULT 59  
AAT76780  
ID AAT76780 standard; DNA; 25 BP.  
XX  
AC AAT76780;  
XX  
DT 15-SEP-1997 (first entry)  
XX  
DE Staphylococcus aureus exfoliative toxin A competitor primer ETA-B2.  
XX  
KW Asymmetric polymerase chain reaction; nucleic acid amplification;  
KW PCR; detection; assay; exfoliative toxin A; ETA; Skin lesion;  
KW competitive primer; capture probe; ss.  
XX  
OS Synthetic.  
XX  
PN US5627054-A.  
XX  
PD 06-MAY-1997.  
XX  
PF 05-APR-1996; 96US-0628417.  
XX  
PR 05-APR-1996; 96US-0628417.  
XX  
PA (USSA) US SEC OF ARMY.  
XX  
PI Gillespie D;  
XX  
DR WPI; 1997-271311/24.  
XX  
PT Quantitative nucleic acid amplification - by competitor primer  
PT asymmetric polymerase chain reaction  
XX  
PS Example 1; Column 5; 9pp; English.  
XX  
CC In a specific example of a novel process for amplifying an amount  
CC (known or unknown) of a double-stranded nucleic acid segment to produce  
CC single-stranded nucleic acid in an amount that is proportional to the  
CC starting amount of the nucleic acid, the Staphylococcus aureus  
CC exfoliative toxin A (ETA) gene was used as the DNA template. The  
CC region comprising nucleotides 165-436 was amplified by symmetric, the  
CC asymmetric or competitor primer asymmetric PCR using the primers  
CC ETA-A2 and ETA-B (see AAT76778 and AAT76779). For asymmetric PCR, the  
CC amount of primer ETA-B was reduced and for competitor primer  
CC asymmetric PCR a competitor primer ETA-B2 (see AAT76780) was added  
CC with upstream primer ETA-A2 after the initial cycling reaction. PCR  
CC products containing ETA-specific sequences were detected  
CC radioactively by a capture system which employed a bifunctional  
CC capture probe ETA-CP (see AAT76781 and AAT76782). ETA-CP was designed  
CC to capture the amplified sense strand onto capture membranes  
CC through hybridisation between the first 40 nucleotides of ETA-CP  
CC and nucleotides 321-360 of the ETA gene and through hybridisation  
CC of the poly(dA) tail on ETA-CP with poly(dT) tails on the capture  
CC membranes. A radioactively labelled "label probe" (see AAT76783),  
CC complementary to nucleotides 389-410 of the ETA gene was used to  
CC detect the amplicons. Results showed that hybridisation of the  
CC capture probe and label probe to the denatured symmetric PCR  
CC product was much less efficient than hybridisation to the  
CC single-stranded PCR products of the asymmetric and competitor  
CC primer asymmetric reactions.  
XX

```
SQ Sequence 25 BP; 4 A; 5 C; 3 G; 8 T; 5 other;
Query Match 100.0%; Score 10; DB 18; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 60
ID ABV82471/c
XX AC ABV82471;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3717.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
XX KW human testis expressed Patched like protein; testis; adrenal; liver;
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00666.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 23-MAY-2001; 2001WO-US00669.
XX PR 09-OCT-2001; 2001US-0864761.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX DR WPI; 2002-676582/73.
XX PT Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 551; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
XX CC has two isoforms, with a few single base pair differences between the
XX CC two. One of the single base pair changes introduces a premature stop
XX CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
XX CC shares an overall structure organisation with the Patched protein. The
XX CC shared structural features strongly imply that HTPL plays a role similar
XX CC to that of Patched, and is a potential tumour suppressor. HTPL is
XX CC important in regulating male germ cell development, and the HTPL gene was
XX CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
XX CC useful for diagnosing a disorder caused by mutation in HTPL, and in
XX CC therapy and manufacture of a medicament for treatment or prevention of
XX CC such disorder associated with decreased expression or activity of human
XX CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
XX CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
XX CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
XX CC clinically useful diagnostic markers and potential therapeutic agents for
XX CC male infertility and cancer. The present oligonucleotide was used in an
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CC example from the invention.
XX SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 61
ID ABV82472/c
XX AC ABV82472;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3718.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
XX KW human testis expressed Patched like protein; testis; adrenal; liver;
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00666.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 23-MAY-2001; 2001WO-US00669.
XX PR 09-OCT-2001; 2001US-0864761.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX DR WPI; 2002-676582/73.
XX PT Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 551; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
XX CC has two isoforms, with a few single base pair differences between the
XX CC two. One of the single base pair changes introduces a premature stop
XX CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
XX CC shares an overall structure organisation with the Patched protein. The
XX CC shared structural features strongly imply that HTPL plays a role similar
XX CC to that of Patched, and is a potential tumour suppressor. HTPL is
XX CC important in regulating male germ cell development, and the HTPL gene was
XX CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
XX CC useful for diagnosing a disorder caused by mutation in HTPL, and in
XX CC therapy and manufacture of a medicament for treatment or prevention of
XX CC such disorder associated with decreased expression or activity of human
XX CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
XX CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
XX CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
XX CC clinically useful diagnostic markers and potential therapeutic agents for
XX CC male infertility and cancer. The present oligonucleotide was used in an
```

CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.  
 XX Sequence 25 BP; 8 A; 2 C; 9 G; 6 T; 0 other;  
 SQ Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 24 CTTCTCTTTT 15

RESULT 62  
 ABV82473/c  
 ID ABV82473 standard; DNA; 25 BP.  
 XX AC ABV82473;  
 XX 03-JAN-2003 (first entry)  
 XX Human HTPL scanning oligonucleotide SEQ ID 3719.  
 XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
 OS Homo sapiens.  
 XX EP1229046-A2.  
 PN 07-AUG-2002.  
 XX 28-JAN-2002; 2002EP-0001167.  
 XX 30-JAN-2001; 2001WO-US00663.  
 PR 30-JAN-2001; 2001WO-US00664.  
 PR 30-JAN-2001; 2001WO-US00665.  
 PR 30-JAN-2001; 2001WO-US00667.  
 PR 30-JAN-2001; 2001WO-US00668.  
 PR 30-JAN-2001; 2001WO-US00669.  
 PR 23-MAY-2001; 2001US-0864761.  
 PR 09-OCT-2001; 2001US-0327898.  
 XX (AEOM-) AEOMICA INC.  
 XX Zhan J;  
 XX WPI; 2002-676582/73.  
 XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX Example 2; Page 551; 718pp; English.

CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and

CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.  
 XX Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;  
 SQ Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 23 CTTCTCTTTT 14

RESULT 63  
 ABV82474/c  
 ID ABV82474 standard; DNA; 25 BP.  
 XX AC ABV82474;  
 XX 03-JAN-2003 (first entry)  
 XX Human HTPL scanning oligonucleotide SEQ ID 3720.  
 XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
 XX Homo sapiens.  
 XX EP1229046-A2.  
 PN 07-AUG-2002.  
 XX 28-JAN-2002; 2002EP-0001167.  
 XX 30-JAN-2001; 2001WO-US00663.  
 PR 30-JAN-2001; 2001WO-US00664.  
 PR 30-JAN-2001; 2001WO-US00665.  
 PR 30-JAN-2001; 2001WO-US00667.  
 PR 30-JAN-2001; 2001WO-US00668.  
 PR 30-JAN-2001; 2001WO-US00669.  
 PR 23-MAY-2001; 2001US-0864761.  
 PR 09-OCT-2001; 2001US-0327898.  
 XX (AEOM-) AEOMICA INC.  
 XX Zhan J;  
 XX WPI; 2002-676582/73.  
 XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX Example 2; Page 551; 718pp; English.

CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and



CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 22 CTTCTCTTTT 13

## RESULT 64

ABV82475/c

ID ABV82475 standard; DNA; 25 BP.

AC ABV82475;

XX 03-JAN-2003 (first entry)

DE Human HTPL scanning oligonucleotide SEQ ID 3721.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

XX EP1229046-A2.

XX 07-AUG-2002.

XX 28-JAN-2002; 2002EP-0001167.

XX 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 30-JAN-2001; 2001WO-US000669.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

PS Example 2; Page 551; 718pp; English.

XX The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are

CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX Sequence 25 BP; 9 A; 2 C; 8 G; 6 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 21 CTTCTCTTTT 12

## RESULT 65

ABV82476/c

ID ABV82476 standard; DNA; 25 BP.

AC ABV82476;

XX 03-JAN-2003 (first entry)

DE Human HTPL scanning oligonucleotide SEQ ID 3722.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

XX EP1229046-A2.

XX 07-AUG-2002.

XX 28-JAN-2002; 2002EP-0001167.

XX 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 30-JAN-2001; 2001WO-US000669.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

PS Example 2; Page 551; 718pp; English.

XX The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is

CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.  
 XX  
 SQ Sequence 25 BP; 9 A; 2 C; 8 G; 6 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 Db 20 CTCTCTCTTT 11  
 |||||

RESULT 66  
 ABV82477/c  
 ID ABV82477 standard; DNA; 25 BP.

XX AC ABV82477;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 3723.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US006663.

XX PR 30-JAN-2001; 2001WO-US006664.

XX PR 30-JAN-2001; 2001WO-US006665.

XX PR 30-JAN-2001; 2001WO-US006667.

XX PR 30-JAN-2001; 2001WO-US006668.

XX PR 23-MAY-2001; 2001WO-US006669.

XX PR 09-OCT-2001; 2001US-0864761.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX  
 PS Example 2; Page 552; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The

CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.  
 XX

SQ Sequence 25 BP; 9 A; 1 C; 8 G; 7 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 Db 19 CTCTCTCTTT 10  
 |||||

RESULT 67  
 ABV82478/c  
 ID ABV82478 standard; DNA; 25 BP.

XX AC ABV82478;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 3724.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US006663.

XX PR 30-JAN-2001; 2001WO-US006664.

XX PR 30-JAN-2001; 2001WO-US006665.

XX PR 30-JAN-2001; 2001WO-US006667.

XX PR 30-JAN-2001; 2001WO-US006668.

XX PR 23-MAY-2001; 2001WO-US006669.

XX PR 09-OCT-2001; 2001US-0864761.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -  
 XX  
 PS Example 2; Page 552; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop

CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
CC shares an overall structure organisation with the Patched protein. The  
CC shared structural features strongly imply that HTPL plays a role similar  
CC to that of Patched, and is a potential tumour suppressor. HTPL is  
CC important in regulating male germ cell development, and the HTPL gene was  
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
CC therapy and manufacture of a medicament for treatment or prevention of  
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
CC clinically useful diagnostic markers and potential therapeutic agents for  
CC male infertility and cancer. The present oligonucleotide was used in an  
CC example from the invention.

XX SQ Sequence 25 BP; 9 A; 1 C; 9 G; 6 T; 0 other;  
Query Match 100.0%; Score 10; DB 24; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 18 CTTCTCTTTT 9

RESULT 68  
ABV82479/c  
ID ABV82479 standard; DNA; 25 BP.  
XX AC ABV82479;  
XX DT 03-JAN-2003 (first entry)  
XX DE Human HTPL scanning oligonucleotide SEQ ID 3725.  
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
XX KW human testis expressed Patched like protein; testis; adrenal; liver;  
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
XX OS Homo sapiens.  
XX PN EPI229046-A2.  
XX PD 07-AUG-2002.  
XX PF 28-JAN-2002; 2002EP-0001167.  
XX PR 30-JAN-2001; 2001WO-US00663.  
XX PR 30-JAN-2001; 2001WO-US00664.  
XX PR 30-JAN-2001; 2001WO-US00665.  
XX PR 30-JAN-2001; 2001WO-US00667.  
XX PR 30-JAN-2001; 2001WO-US00668.  
XX PR 30-JAN-2001; 2001WO-US00669.  
XX PR 23-MAY-2001; 2001US-0864761.  
XX PR 09-OCT-2001; 2001US-0327898.  
XX PA (AEOM-) AEOMICA INC.  
XX PI Zhan J;  
XX WPI; 2002-676582/73.  
XX DR Novel isolated human testis expressed Patched like protein (HTPL),  
XX PT useful for identifying agonist and antagonist and specific binding  
XX PT partners, and for treating subjects having defects in HTPL -  
XX PS Example 2; Page 552; 718pp; English.  
XX CC The present invention relates to human testis expressed Patched like  
XX protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL

CC has two isoforms, with a few single base pair differences between the  
CC two. One of the single base pair changes introduces a premature stop  
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
CC shares an overall structure organisation with the Patched protein. The  
CC shared structural features strongly imply that HTPL plays a role similar  
CC to that of Patched, and is a potential tumour suppressor. HTPL is  
CC important in regulating male germ cell development, and the HTPL gene was  
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
CC therapy and manufacture of a medicament for treatment or prevention of  
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
CC clinically useful diagnostic markers and potential therapeutic agents for  
CC male infertility and cancer. The present oligonucleotide was used in an  
CC example from the invention.

XX SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;  
Query Match 100.0%; Score 10; DB 24; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 17 CTTCTCTTTT 8

RESULT 69  
ABV82480/c  
ID ABV82480 standard; DNA; 25 BP.  
XX AC ABV82480;  
XX DT 03-JAN-2003 (first entry)  
XX DE Human HTPL scanning oligonucleotide SEQ ID 3726.  
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
XX KW human testis expressed Patched like protein; testis; adrenal; liver;  
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
XX OS Homo sapiens.  
XX PN EPI229046-A2.  
XX PD 07-AUG-2002.  
XX PF 28-JAN-2002; 2002EP-0001167.  
XX PR 30-JAN-2001; 2001WO-US00663.  
XX PR 30-JAN-2001; 2001WO-US00664.  
XX PR 30-JAN-2001; 2001WO-US00665.  
XX PR 30-JAN-2001; 2001WO-US00667.  
XX PR 30-JAN-2001; 2001WO-US00668.  
XX PR 30-JAN-2001; 2001WO-US00669.  
XX PR 23-MAY-2001; 2001US-0864761.  
XX PR 09-OCT-2001; 2001US-0327898.  
XX PA (AEOM-) AEOMICA INC.  
XX PI Zhan J;  
XX WPI; 2002-676582/73.  
XX DR Novel isolated human testis expressed Patched like protein (HTPL),  
XX PT useful for identifying agonist and antagonist and specific binding  
XX PT partners, and for treating subjects having defects in HTPL -  
XX PS Example 2; Page 552; 718pp; English.  
XX CC The present invention relates to human testis expressed Patched like  
XX protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL

CC The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

XX  
 SQ Sequence 25 BP; 9 A; 3 C; 8 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10  
 |||||  
 Db 16 CTCTCTCTTT 7

RESULT 70  
 ABV82481/c  
 ID ABV82481 standard; DNA; 25 BP.

AC ABV82481;

XX 03-JAN-2003 (first entry)

XX Human HTPL scanning oligonucleotide SEQ ID 3727.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

XX EP1229046-A2.

XX 07-AUG-2002.

XX 28-JAN-2002; 2002EP-0001167.

XX 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding  
 PT partners, and for treating subjects having defects in HTPL -

XX

PS Example 2; Page 552; 718pp; English.

XX The present invention relates to human testis expressed Patched like  
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
 CC has two isoforms, with a few single base pair differences between the  
 CC two. One of the single base pair changes introduces a premature stop  
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
 CC shares an overall structure organisation with the Patched protein. The  
 CC shared structural features strongly imply that HTPL plays a role similar  
 CC to that of Patched, and is a potential tumour suppressor. HTPL is  
 CC important in regulating male germ cell development, and the HTPL gene was  
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
 CC therapy and manufacture of a medicament for treatment or prevention of  
 CC such disorder associated with decreased expression or activity of human  
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
 CC clinically useful diagnostic markers and potential therapeutic agents for  
 CC male infertility and cancer. The present oligonucleotide was used in an  
 CC example from the invention.

SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

|||||  
 Db 15 CTCTCTCTTT 6

RESULT 71

ABV82482/c

ID ABV82482 standard; DNA; 25 BP.

XX ABV82482;

XX 03-JAN-2003 (first entry)

XX Human HTPL scanning oligonucleotide SEQ ID 3728.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
 KW human testis expressed Patched like protein; testis; adrenal; liver;  
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

XX EP1229046-A2.

XX 07-AUG-2002.

XX 28-JAN-2002; 2002EP-0001167.

XX 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),  
 PT useful for identifying agonist and antagonist and specific binding

PT useful for identifying agonist and antagonist and specific binding

PT partners, and for treating subjects having defects in HTPL -  
XX  
PS Example 2; Page 552; 718pp; English.  
XX  
CC The present invention relates to human testis expressed Patched like  
CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
CC has two isoforms, with a few single base pair differences between the  
CC two. One of the single base pair changes introduces a premature stop  
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
CC shares an overall structure organisation with the Patched protein. The  
CC shared structural features strongly imply that HTPL plays a role similar  
CC to that of Patched, and is a potential tumour suppressor. HTPL is  
CC important in regulating male germ cell development, and the HTPL gene was  
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
CC therapy and manufacture of a medicament for treatment or prevention of  
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
CC clinically useful diagnostic markers and potential therapeutic agents for  
CC male infertility and cancer. The present oligonucleotide was used in an  
CC example from the invention.  
SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;  
Query Match 100.0%; Score 10; DB 24; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 14 CTTCTCTTTT 5  
RESULT 72  
ABV82483/c  
ID ABV82483 standard; DNA; 25 BP.  
XX  
AC ABV82483;  
XX  
DT 03-JAN-2003 (first entry)  
XX  
DE Human HTPL scanning oligonucleotide SEQ ID 3729.  
XX  
KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
KW human testis expressed Patched like protein; testis; adrenal; liver;  
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
XX  
OS Homo sapiens.  
XX  
PN EPI229046-A2.  
XX  
PD 07-AUG-2002.  
XX  
PF 28-JAN-2002; 2002EP-0001167.  
XX  
PR 30-JAN-2001; 2001WO-US00663.  
XX  
PR 30-JAN-2001; 2001WO-US00664.  
XX  
PR 30-JAN-2001; 2001WO-US00665.  
XX  
PR 30-JAN-2001; 2001WO-US00667.  
XX  
PR 30-JAN-2001; 2001WO-US00668.  
XX  
PR 30-JAN-2001; 2001WO-US00669.  
XX  
PR 23-MAY-2001; 2001US-0864761.  
XX  
PR 09-OCT-2001; 2001US-0327898.  
XX  
FA (AEOM-) AEOMICA INC.  
XX  
PI Zhan J;  
XX  
DR WPI; 2002-676582/73.  
XX

PT Novel isolated human testis expressed Patched like protein (HTPL),  
XX useful for identifying agonist and antagonist and specific binding  
XX partners, and for treating subjects having defects in HTPL -  
XX Example 2; Page 552; 718pp; English.  
XX  
CC The present invention relates to human testis expressed Patched like  
CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL  
CC has two isoforms, with a few single base pair differences between the  
CC two. One of the single base pair changes introduces a premature stop  
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL  
CC shares an overall structure organisation with the Patched protein. The  
CC shared structural features strongly imply that HTPL plays a role similar  
CC to that of Patched, and is a potential tumour suppressor. HTPL is  
CC important in regulating male germ cell development, and the HTPL gene was  
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are  
CC useful for diagnosing a disorder caused by mutation in HTPL, and in  
CC therapy and manufacture of a medicament for treatment or prevention of  
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and  
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,  
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are  
CC clinically useful diagnostic markers and potential therapeutic agents for  
CC male infertility and cancer. The present oligonucleotide was used in an  
CC example from the invention.  
SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;  
Query Match 100.0%; Score 10; DB 24; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 13 CTTCTCTTTT 4  
RESULT 73  
ABV82484/c  
ID ABV82484 standard; DNA; 25 BP.  
XX  
AC ABV82484;  
XX  
DT 03-JAN-2003 (first entry)  
XX  
DE Human HTPL scanning oligonucleotide SEQ ID 3730.  
XX  
KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;  
KW human testis expressed Patched like protein; testis; adrenal; liver;  
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;  
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.  
XX  
OS Homo sapiens.  
XX  
PN EPI229046-A2.  
XX  
PD 07-AUG-2002.  
XX  
PF 28-JAN-2002; 2002EP-0001167.  
XX  
PR 30-JAN-2001; 2001WO-US00663.  
XX  
PR 30-JAN-2001; 2001WO-US00664.  
XX  
PR 30-JAN-2001; 2001WO-US00665.  
XX  
PR 30-JAN-2001; 2001WO-US00667.  
XX  
PR 30-JAN-2001; 2001WO-US00668.  
XX  
PR 30-JAN-2001; 2001WO-US00669.  
XX  
PR 23-MAY-2001; 2001US-0864761.  
XX  
PR 09-OCT-2001; 2001US-0327898.  
XX  
FA (AEOM-) AEOMICA INC.  
XX  
PI Zhan J;  
XX  
DR WPI; 2002-676582/73.  
XX

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DR  WPI; 2002-676582/73.
XX
PT  Novel isolated human testis expressed Patched like protein (HTPL),
PT  useful for identifying agonist and antagonist and specific binding
XX  partners, and for treating subjects having defects in HTPL -
PS  Example 2; Page 552; 718pp; English.
XX
CC  The present invention relates to human testis expressed Patched like
CC  protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC  has two isoforms, with a few single base pair differences between the
CC  two. One of the single base pair changes introduces a premature stop
CC  codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC  shares an overall structure organisation with the Patched protein. The
CC  shared structural features strongly imply that HTPL plays a role similar
CC  to that of Patched, and is a potential tumour suppressor. HTPL is
CC  important in regulating male germ cell development, and the HTPL gene was
CC  mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC  useful for diagnosing a disorder caused by mutation in HTPL, and in
CC  therapy and manufacture of a medicament for treatment or prevention of
CC  such disorder associated with decreased expression or activity of human
CC  HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC  foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC  skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC  clinically useful diagnostic markers and potential therapeutic agents for
CC  male infertility and cancer. The present oligonucleotide was used in an
CC  example from the invention.
XX  SQ  Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;
XX
XX  Query Match 100.0%; Score 10; DB 24; Length 25;
XX  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY  1 CTTCTCTTTT 10
Db  12 CTTCTCTTTT 3
XX
RESULT 74
ABV82485/c
ID  ABV82485 standard; DNA; 25 BP.
AC  ABV82485;
XX
DT  03-JAN-2003 (first entry)
XX
DE  Human HTPL scanning oligonucleotide SEQ ID 3731.
XX
KW  Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW  human testis expressed Patched like protein; testis; adrenal; liver;
KW  male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW  prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
OS  Homo sapiens.
XX
PN  EP1229046-A2.
XX
PD  07-AUG-2002.
XX
PF  28-JAN-2002; 2002EP-0001167.
XX
PR  30-JAN-2001; 2001WO-US00663.
PR  30-JAN-2001; 2001WO-US00664.
PR  30-JAN-2001; 2001WO-US00665.
PR  30-JAN-2001; 2001WO-US00667.
PR  30-JAN-2001; 2001WO-US00668.
PR  30-JAN-2001; 2001WO-US00669.
PR  23-MAY-2001; 2001US-0864761.
PR  09-OCT-2001; 2001US-0327898.
XX
PA  (AEOM-) AEOMICA INC.
XX
PI  Zhan J;
XX
DR  WPI; 2002-676582/73.
XX
PT  Novel isolated human testis expressed Patched like protein (HTPL),
PT  useful for identifying agonist and antagonist and specific binding
XX  partners, and for treating subjects having defects in HTPL -
PS  Example 2; Page 553; 718pp; English.
XX
CC  The present invention relates to human testis expressed Patched like
CC  protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC  has two isoforms, with a few single base pair differences between the
CC  two. One of the single base pair changes introduces a premature stop
CC  codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC  shares an overall structure organisation with the Patched protein. The
CC  shared structural features strongly imply that HTPL plays a role similar
CC  to that of Patched, and is a potential tumour suppressor. HTPL is
CC  important in regulating male germ cell development, and the HTPL gene was
CC  mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC  useful for diagnosing a disorder caused by mutation in HTPL, and in
CC  therapy and manufacture of a medicament for treatment or prevention of
CC  such disorder associated with decreased expression or activity of human
CC  HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC  foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC  skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC  clinically useful diagnostic markers and potential therapeutic agents for
CC  male infertility and cancer. The present oligonucleotide was used in an
CC  example from the invention.
XX  SQ  Sequence 25 BP; 9 A; 4 C; 6 G; 6 T; 0 other;
XX
XX  Query Match 100.0%; Score 10; DB 24; Length 25;
XX  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY  1 CTTCTCTTTT 10
Db  11 CTTCTCTTTT 2
XX
RESULT 75
ABV82486/c
ID  ABV82486 standard; DNA; 25 BP.
AC  ABV82486;
XX
DT  03-JAN-2003 (first entry)
XX
DE  Human HTPL scanning oligonucleotide SEQ ID 3732.
XX
KW  Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW  human testis expressed Patched like protein; testis; adrenal; liver;
KW  male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW  prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
OS  Homo sapiens.
XX
PN  EP1229046-A2.
XX
PD  07-AUG-2002.
XX
PF  28-JAN-2002; 2002EP-0001167.
XX
PR  30-JAN-2001; 2001WO-US00663.
PR  30-JAN-2001; 2001WO-US00664.
PR  30-JAN-2001; 2001WO-US00665.
PR  30-JAN-2001; 2001WO-US00667.
PR  30-JAN-2001; 2001WO-US00668.
PR  30-JAN-2001; 2001WO-US00669.
PR  23-MAY-2001; 2001US-0864761.
PR  09-OCT-2001; 2001US-0327898.
XX

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PA (AEOM-) AEOMICA INC.
XX
PI Zhan J;
XX
XX WPI; 2002-676582/73.
XX
XX Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
XX Example 2; Page 553; 718pp; English.
XX
XX The present invention relates to human testis expressed Patched like
CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC such disorder associated with decreased expression or activity of human
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
XX Sequence 25 BP; 9 A; 4 C; 6 G; 6 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1
XX
RESULT 76
ABS75576/c
ID ABS75576 standard; DNA; 25 BP.
XX
AC ABS75576;
XX
XX 27-DEC-2002 (first entry)
XX
XX Human PAPP-Ea associated 25-mer SEQ ID 1102.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX dysgenetic pregnancy; primer; ss.
XX
XX Homo sapiens.
XX
XX OS
XX US2002102252-A1.
XX
XX 01-AUG-2002.
XX
XX 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX (GUYY/) GU Y.
XX (SHAN/) SHANNON M E.
XX
XX Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
XX associated plasma protein E, for preventing or aborting pregnancy -
XX
XX Example 2; Page 220; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or
CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
CC antibodies can be used to assess the expression levels of PAPP-E isoform
CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
CC antenatally. This sequence represents an oligomer used in scanning the
CC human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 11 A; 3 C; 8 G; 3 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16
XX
RESULT 77
ABS75577/c
ID ABS75577 standard; DNA; 25 BP.
XX
AC ABS75577;
XX
XX 27-DEC-2002 (first entry)
XX
XX Human PAPP-Ea associated 25-mer SEQ ID 1103.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX dysgenetic pregnancy; primer; ss.
XX
XX Homo sapiens.
XX
XX OS
XX US2002102252-A1.
XX
XX 01-AUG-2002.
XX
XX 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX (GUYY/) GU Y.
XX (SHAN/) SHANNON M E.
XX
XX Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
XX associated plasma protein E, for preventing or aborting pregnancy -
XX
XX Example 2; Page 220; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or

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CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 11 A; 3 C; 8 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 24 CTTCTCTTTT 15

RESULT 78  
 ABS75578/c  
 ID ABS75578 standard; DNA; 25 BP.  
 AC ABS75578;  
 XX  
 XX 27-DEC-2002 (first entry)  
 DT Human PAPP-Ea associated 25-mer SEQ ID 1104.  
 XX  
 XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX Homo sapiens.  
 OS  
 XX US2002102252-A1.  
 PN  
 XX 01-AUG-2002.  
 PD  
 XX 06-APR-2001; 2001US-0827998.  
 PF  
 XX 26-MAY-2000; 2000US-207456P.  
 PR (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX Gu Y, Shannon ME;  
 PI WPI; 2002-697817/75.  
 DR  
 XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy -  
 PF Example 2; Page 220; 353pp; English.  
 PS  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 11 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 24 CTTCTCTTTT 15

RESULT 80  
 ABS75580/c  
 ID ABS75580 standard; DNA; 25 BP.  
 AC ABS75580;  
 XX  
 XX 27-DEC-2002 (first entry)  
 DT Human PAPP-Ea associated 25-mer SEQ ID 1105.  
 XX  
 XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX Homo sapiens.  
 OS  
 XX US2002102252-A1.  
 PN  
 XX 01-AUG-2002.  
 PD  
 XX 06-APR-2001; 2001US-0827998.  
 PF  
 XX 26-MAY-2000; 2000US-207456P.  
 PR (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX Gu Y, Shannon ME;  
 PI WPI; 2002-697817/75.  
 DR  
 XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy -  
 PF Example 2; Page 220; 353pp; English.  
 PS  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 11 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 22 CTTCTCTTTT 13

RESULT 80  
 ABS75580/c  
 ID ABS75580 standard; DNA; 25 BP.  
 AC ABS75580;  
 XX  
 XX 27-DEC-2002 (first entry)  
 DT Human PAPP-Ea associated 25-mer SEQ ID 1105.  
 XX  
 XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX Homo sapiens.  
 OS  
 XX US2002102252-A1.  
 PN  
 XX 01-AUG-2002.  
 PD  
 XX 06-APR-2001; 2001US-0827998.  
 PF  
 XX 26-MAY-2000; 2000US-207456P.  
 PR (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX Gu Y, Shannon ME;  
 PI WPI; 2002-697817/75.  
 DR  
 XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy -  
 PF Example 2; Page 220; 353pp; English.  
 PS  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 22 CTTCTCTTTT 13

RESULT 80  
 ABS75580/c  
 ID ABS75580 standard; DNA; 25 BP.  
 AC ABS75580;





DR WPI; 2002-697817/75.  
 XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 PS Example 2; Page 221; 353pp; English.  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Qy 1 CTTCTCTTTT 10  
 Db 19 CTTCTCTTTT 10  
 RESULT 83  
 ABS75583/C  
 ID ABS75583 standard; DNA; 25 BP.  
 AC ABS75583;  
 XX  
 DT 27-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 25-mer SEQ ID 1109.  
 XX  
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN US2002102252-A1.  
 XX  
 PD 01-AUG-2002.  
 XX  
 PF 06-APR-2001; 2001US-0827998.  
 XX  
 PR 26-MAY-2000; 2000US-207456P.  
 XX  
 PA (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX  
 PI Gu Y, Shannon ME;  
 XX  
 DR WPI; 2002-697817/75.  
 XX  
 PT New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 PS Example 2; Page 221; 353pp; English.  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be

CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 12 A; 3 C; 6 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Qy 1 CTTCTCTTTT 10  
 Db 18 CTTCTCTTTT 9  
 RESULT 84  
 ABS75584/C  
 ID ABS75584 standard; DNA; 25 BP.  
 AC ABS75584;  
 XX  
 DT 27-DEC-2002 (first entry)  
 XX  
 DE Human PAPP-Ea associated 25-mer SEQ ID 1110.  
 XX  
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN US2002102252-A1.  
 XX  
 PD 01-AUG-2002.  
 XX  
 PF 06-APR-2001; 2001US-0827998.  
 XX  
 PR 26-MAY-2000; 2000US-207456P.  
 XX  
 PA (GUY/) GU Y.  
 PA (SHAN/) SHANNON M E.  
 XX  
 PI Gu Y, Shannon ME;  
 XX  
 DR WPI; 2002-697817/75.  
 XX  
 PT New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy  
 PS Example 2; Page 221; 353pp; English.  
 XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 12 A; 4 C; 5 G; 4 T; 0 other;  
 Query Match 100.0%; Score 10; DB 24; Length 25;

```

Best Local Similarity 100.0%; Pred. NO. 1.8e+04; Indels 0; Gaps 0;
Matches 10; Conservative 0; Mismatches 0;

Qy 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 85
ABS75585/c
ID ABS75585 standard; DNA; 25 BP.
XX
AC ABS75585;
XX
DT 27-DEC-2002 (first entry)
XX
DE Human PAPP-Ea associated 25-mer SEQ ID 1111.
XX
KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
dysgenetic pregnancy; primer; ss.
XX
OS Homo sapiens.
XX
PN US2002102252-A1.
XX
PD 01-AUG-2002.
XX
PF 06-APR-2001; 2001US-0827998.
XX
PR 26-MAY-2000; 2000US-207456P.
XX
PA (GUY/) GU Y.
PA (SHAN/) SHANNON M E.
XX
PI Gu Y, Shannon ME;
XX
PS WPI; 2002-697817/75.
XX
PT New isolated nucleic acid encoding an isoform of human pregnancy
associated plasma protein E, for preventing or aborting pregnancy -
XX
PS Example 2; Page 221; 353pp; English.
XX
CC This invention describes a novel isolated nucleic acid that encodes
one of three new isoforms of human pregnancy associated plasma protein E,
hPAPP-E. The products of the invention have abortive and contraceptive
activity and can be used for gene therapy or in a vaccine. The nucleic
acid, polypeptide encoded by it, or antibody to the polypeptide can be
used in pharmaceutical compositions or vaccines for preventing or
aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
dysgenetic pregnancies. The nucleic acids are used as probes to assess
the level of PAPP-E isoform mRNA in chorionic villus samples, and the
antibodies can be used to assess the expression levels of PAPP-E isoform
proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
antenatally. This sequence represents an oligomer used in scanning the
human PAPP-E genes described in the disclosure of the invention.
XX
SQ Sequence 25 BP; 12 A; 4 C; 4 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 16 CTTCTCTTTT 7

RESULT 86
ABS75586/c
ID ABS75586 standard; DNA; 25 BP.
XX

```



CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.  
 XX  
 SQ Sequence 25 BP; 11 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10  
 |||||  
 Db 12 CTCTCTCTTTT 3

RESULT 90  
 ABS75590/c  
 ID ABS75590 standard; DNA; 25 BP.

XX AC ABS75590;

XX DT 27-DEC-2002 (first entry)

DE Human PAPP-Ea associated 25-mer SEQ ID 1116.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.

XX OS Homo sapiens.

XX PN US2002102252-A1.

XX PD 01-AUG-2002.

XX PF 06-APR-2001; 2001US-0827998.

XX PR 26-MAY-2000; 2000US-207456P.

XX PA (GUY/) GU Y.

XX PA (SHAN/) SHANNON M E.

XX PI Gu Y, Shannon ME;

XX PR WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy

XX Example 2; Page 222; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.

XX Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10  
 |||||  
 Db 11 CTCTCTCTTTT 2

RESULT 91

ABS75591/c  
 ID ABS75591 standard; DNA; 25 BP.

XX AC ABS75591;

XX DT 27-DEC-2002 (first entry)

XX Human PAPP-Ea associated 25-mer SEQ ID 1117.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;  
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;  
 KW dysgenetic pregnancy; primer; ss.

XX OS Homo sapiens.

XX PN US2002102252-A1.

XX PD 01-AUG-2002.

XX PF 06-APR-2001; 2001US-0827998.

XX PR 26-MAY-2000; 2000US-207456P.

XX PA (GUY/) GU Y.

XX PA (SHAN/) SHANNON M E.

XX PI Gu Y, Shannon ME;

XX PR WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy  
 PT associated plasma protein E, for preventing or aborting pregnancy

XX Example 2; Page 222; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes  
 CC one of three new isoforms of human pregnancy associated plasma protein E,  
 CC hPAPP-E. The products of the invention have abortive and contraceptive  
 CC activity and can be used for gene therapy or in a vaccine. The nucleic  
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be  
 CC used in pharmaceutical compositions or vaccines for preventing or  
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of  
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess  
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the  
 CC antibodies can be used to assess the expression levels of PAPP-E isoform  
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies  
 CC antenatally. This sequence represents an oligomer used in scanning the  
 CC human PAPP-E genes described in the disclosure of the invention.

XX Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10  
 |||||  
 Db 10 CTCTCTCTTTT 1

RESULT 92

AAQ45398/c  
 ID AAQ45398 standard; DNA; 27 BP.

XX AC AAQ45398;  
 XX XX  
 DT 25-MAR-2003 (updated)  
 DT 11-NOV-1994 (first entry)  
 XX XX  
 DE Oligonucleotide forming triplex with viral polypurine tract.  
 XX XX  
 KW HIV; human immunodeficiency virus; retrovirus; hepatitis virus;  
 KW reverse transcription; virus replication; inhibition; treatment;  
 KW therapy; polypurine; triplex; antisense; ss.  
 XX OS Synthetic.  
 XX XX  
 FN WO9407367-A1.  
 XX PD 14-APR-1994.  
 XX XX  
 PF 29-SEP-1993; 93WO-US09300.  
 XX XX  
 PR 29-SEP-1992; 92US-0954184.  
 XX XX  
 PA (APOL-) APOLLON INC.  
 PA (PLAC ) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.  
 XX XX  
 PI Moelling K;  
 XX XX  
 DR WPI; 1994-135099/16.  
 XX XX  
 XX Antiviral oligomers that bind poly-purine tracts of  
 PT single-stranded RNA or RNA - DNA hybrids - used to target the  
 PT early stages of viral replication before double stranded DNA is  
 PT formed  
 XX XX  
 PS Example 4; Page 29; 54pp; English.  
 XX XX  
 CC Administration of antisense or triplex forming oligonucleotides  
 CC which bind polypurine tracts (PPT) may be used in the therapy or  
 CC treatment of individuals infected with retroviruses or hepatitis  
 CC viruses since in these two families of viruses, two primers are  
 CC involved in the reverse transcription of viral RNA into double  
 CC stranded DNA, one of which is a PPT. The antisense or triplex  
 CC forming oligonucleotides can inhibit the early stages of viral  
 CC replication by binding to the PPT primer or by binding to PPT  
 CC tracts in the RNA-DNA hybrid molecule formed after reverse  
 CC transcription of the viral RNA. This oligonucleotide was incubated  
 CC with an in vitro transcribed 5' end labelled pKJ2 RNA of 134  
 CC nucleotides in length to which a 40-mer deoxyribonucleotide  
 CC complementary to the PPT had been hybridised. The presence of this  
 CC sequence led to protection of the pKJ2 RNA-DNA hybrid from RNase H  
 CC digestion which would suggest the formation of a triplex. Triplex  
 CC formation was confirmed using a primer extension technique. A  
 CC primer binding downstream of the PPT was synthesised and extended in  
 CC vitro by reverse transcriptase in the presence of  
 CC oligodeoxynucleotides including one which was radioactively labelled.  
 CC The newly synthesised DNA was terminated at the site of the PPT when  
 CC the triplex was formed and blocked extension. Triplex formation  
 CC would be expected to interfere with viral replication in vivo.  
 CC See AAQ45381-Q45417. This is a variant of the sequence described in  
 CC AAQ45387.  
 CC (Updated on 25-MAR-2003 to correct PN field.)  
 XX XX  
 SQ Sequence 27 BP; 7 A; 8 C; 8 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 15; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 DB 22 CTTCTCTTTT 13

RESULT 93  
 AAX74299/C  
 ID AAX74299 standard; RNA; 27 BP.  
 XX AC AAX74299;  
 XX XX  
 DT 28-JUL-1999 (first entry)  
 XX XX  
 DE Mouse flt-1 VEGF receptor hammerhead ribozyme #771.  
 XX XX  
 KW Vascular endothelial growth factor receptor; VEGF receptor; flt-1;  
 KW flk-1; KDR; hammerhead ribozyme; hairpin ribozyme; cleavage;  
 KW tumour angiogenesis; psoriasis; rheumatoid arthritis; ocular disease;  
 KW fms-like tyrosine kinase 1; kinase insert domain containing receptor;  
 KW foetal liver kinase 1; ss.  
 XX OS Synthetic.  
 OS Mus sp.  
 XX XX  
 FN WO9715662-A2.  
 XX XX  
 PD 01-MAY-1997.  
 XX XX  
 PF 25-OCT-1996; 96WO-US17480.  
 XX XX  
 PR 11-JAN-1996; 96US-0584040.  
 PR 26-OCT-1995; 95US-0005974.  
 XX XX  
 PA (CHIR ) CHIRON CORP.  
 PA (RIBO-) RIBOZYME PHARM INC.  
 XX XX  
 PI Escobedo J, McSwiggen J, Pavco P, Stinchcomb D;  
 XX XX  
 DR WPI; 1997-259017/23.  
 XX XX  
 PT Nucleic acid molecule modulating VEGF receptor(s) gene expression or  
 PT mRNA stability - useful for treating e.g. tumour angiogenesis,  
 PT psoriasis, rheumatoid arthritis, etc., in a human patient  
 XX XX  
 PS Claim 9; Page 178; 218pp; English.  
 XX CC  
 CC The present invention describes nucleic acid molecules which modulate  
 CC the synthesis, expression and/or stability of a mRNA encoding 1 or more  
 CC receptors of vascular endothelial growth factor (VEGF). A patient  
 CC (preferably human) having a condition associated with the level of the  
 CC fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing  
 CC receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour  
 CC angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can  
 CC be treated by administering the nucleic acid molecule or the expression  
 CC vector to the patient. AAX67275 to AAX75752 represent specific examples  
 CC of nucleic acid molecules from the present invention.  
 XX XX  
 SQ Sequence 27 BP; 12 A; 1 C; 9 G; 4 U; 1 other;

Query Match 100.0%; Score 10; DB 18; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 DB 27 CTTCTCTTTT 18

RESULT 94  
 AAH62997  
 ID AAH62997 standard; DNA; 27 BP.  
 XX AC AAH62997;  
 XX XX  
 DT 11-SEP-2001 (first entry)  
 XX XX  
 DE Shrimp white spot Bacilliform virus (WSBV) oligonucleotide 158.  
 XX XX

KW Shrimp white spot Bacilliform virus; WSBV; diagnosis; viral infection;  
KW antiviral agent; gene expression; antisense construct; probe; primer;  
KW transgenic viral resistant shrimp; ss.  
XX  
XX  
OS White spot syndrome virus.  
XX  
XX WO200138351-A2.  
XX  
XX 31-MAY-2001.  
XX  
XX 08-NOV-2000; 2000WO-US28888.  
XX  
XX 24-NOV-1999; 99CN-0124717.  
XX  
XX (PENY-) PE CORP NY.  
PA  
PA (THIR-) THIRD INST OCEANOGRAPHY STATE OCEANI C A.  
PA (SINO-) SINOGENOMAX CO LTD.  
XX  
XX Xu X, Yang F, He J, Pham L, He M, Ye Y, Shen Y, Kodira C;  
XX  
XX WPI; 2001-355877/37.  
XX  
XX Primary nucleotide sequence of the shrimp white spot Bacilliform virus  
PT (WSBV), useful for producing viral polypeptides that can be used to  
PT screen for agents that are useful for treating WSBV infection -  
XX  
XX Disclosure; Figure 3; 626pp; English.  
XX  
XX The invention provides the primary nucleotide sequence of the WSBV genome  
CC (AAH62689), predicted transcript sequences (AAH62689-AAH62839) and  
CC encoded proteins (AAH62840-AAH62851) and oligonucleotide sequences  
CC (AAH62840-63160) suitable for use as primers or probes. The nucleic acid  
CC molecules and proteins of the invention are useful for diagnosis and  
CC monitoring viral infection, in screens for antiviral agents and for  
CC monitoring viral gene expression or activity during a treatment regimen.  
CC The nucleic acid molecules are also useful as antisense constructs to  
CC control viral gene expression in infected cells and tissues and to create  
CC transgenic viral resistant shrimp.  
XX  
SQ Sequence 27 BP; 5 A; 6 C; 2 G; 14 T; 0 other;  
  
Query Match 100.0%; Score 10; DB 22; Length 27;  
Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
Qy 1 CTCTCTCTTT 10  
Db 2 CTCTCTCTTT 11  
  
RESULT 95  
AAH38166/C  
ID AAH38166 standard; DNA; 27 BP.  
XX  
XX AAH38166;  
XX  
XX 14-AUG-2001 (first entry)  
XX  
XX SNP specific lower PCR primer SEQ ID 962.  
XX  
XX Single nucleotide polymorphism; SNP; single nucleotide primer extension;  
KW SNPE; genotyping; agammaglobulinaemia; diabetes insipidus; cancer;  
KW Lesch-Nyhan syndrome; muscular dystrophy; familial hypercholesterolaemia;  
KW polycystic kidney disease; osteogenesis imperfecta; autoimmune disease;  
KW acute intermittent porphyria; rheumatoid arthritis; multiple sclerosis;  
KW inflammation; forensic investigation; paternity analysis; PCR primer; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO200129262-A2.  
PN  
XX  
XX 26-APR-2001.  
PD  
XX

PF 13-OCT-2000; 2000WO-US28436.  
XX  
XX 15-OCT-1999; 99US-0160096.  
XX  
XX (ORCH-) ORCHID BIOSCIENCES INC.  
PA  
XX Picoult-Newburg L, Pohl M;  
PI  
XX WPI; 2001-290930/30.  
DR  
XX New genotyping oligonucleotide, useful for detecting the presence,  
PT absence or identity of single polynucleotide polymorphism in a nucleic  
PT acid sample -  
XX  
XX Claim 1; Page 54; 83pp; English.  
PS  
XX Sequences AAH37205 - AAH40944 represent PCR primers, single nucleotide  
CC primer extension (SNPE) primers, and the sequences of regions flanking  
CC sites of single nucleotide polymorphisms SNPs. The present invention  
CC includes kits for determining the presence or absence of a SNP, using the  
CC oligonucleotides of the invention. The PCR primers are used to amplify a  
CC SNP flanking sequence, the SNPE primer is used as a genotyping primer.  
CC The oligonucleotides are useful for genotyping a nucleic acid sample by  
CC performing a single-nucleotide primer extension reaction. The  
CC oligonucleotides are useful for determining the presence, absence or  
CC identity of a SNP and for genotyping nucleic acid samples, for e.g. to  
CC assess by association analysis the genotype of an individual or group of  
CC individuals, having a pathological phenotypic trait suspected of being  
CC caused by one or more SNPs. Phenotypic traits include diseases e.g.  
CC agammaglobulinaemia, diabetes insipidus, Lesch-Nyhan syndrome, muscular  
CC dystrophy, familial hypercholesterolaemia, polycystic kidney disease,  
CC osteogenesis imperfecta and acute intermittent porphyria. Phenotypic  
CC traits also include symptoms of or susceptibility to multifactorial  
CC disease of which a component is or may be genetic such as autoimmune  
CC diseases, including, rheumatoid arthritis, multiple sclerosis,  
CC inflammation, cancer, nervous system diseases and infection by pathogenic  
CC microorganism. The method is also useful in forensic investigations and  
CC paternity analysis. The present sequence represents a PCR primer specific  
CC for a human SNP containing DNA sequence.  
XX  
SQ Sequence 27 BP; 13 A; 5 C; 4 G; 5 T; 0 other;  
  
Query Match 100.0%; Score 10; DB 22; Length 27;  
Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
Qy 1 CTCTCTCTTT 10  
Db 21 CTCTCTCTTT 12  
  
RESULT 96  
AAC81508  
ID AAC81508 standard; DNA; 28 BP.  
XX  
XX AAC81508;  
AC  
XX 28-FEB-2001 (first entry)  
DT  
XX Human MSF exon 6 RT-PCR primer, SEQ ID NO:16.  
DE  
XX Human MSF; megakaryocyte stimulating factor; tribonectin;  
KW alternative splicing; joint boundary lubricant; O-linked oligosaccharide;  
KW osteoarthritis; tribosupplementation; tissue adhesion inhibition;  
KW friction coefficient reduction; gene therapy; antiarthritic;  
KW osteopathic; reverse transcription-PCR; RT-PCR primer; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO200064930-A2.  
PN  
XX  
XX 02-NOV-2000.  
PD  
XX

PF 24-APR-2000; 2000WO-US10953.  
XX  
XX  
PR 23-APR-1999; 99US-0298970.  
XX  
XX (RHOD-) RHODE ISLAND HOSPITAL LIFESPAN PARTNER.  
PA Jay GD;  
PI WPI; 2001-024673/03.  
XX  
XX Novel tribonectin polypeptide useful as lubricant for treating  
PT osteoarthritis, comprises O-linked lubricating moiety -  
XX Example 1; Page 23; 47pp; English.  
XX  
XX The invention relates to a human tribonectin which is a product of  
CC alternative splicing of the human MSF (megakaryocyte stimulating factor)  
CC gene. The tribonectin has at least one O-linked oligosaccharide  
CC lubricating moiety and has a polypeptide sequence comprising 1-76  
CC repeats of a motif having at least 50% identity to the sequence KEPAPTT  
CC (AA829774). The invention also relates to a nucleic acid encoding a  
CC human MSF-derived tribonectin; a biocompatible composition comprising a  
CC human tribonectin for inhibiting tissue adhesion formation; and a method  
CC of diagnosing osteoarthritis or a predisposition to osteoarthritis by  
CC measuring the amount of MSF or its fragment in a biological sample of a  
CC mammal, wherein an increased amount of MSF compared to a control  
CC indicates the presence of or predisposition to developing  
CC osteoarthritis. The tribonectin and DNA encoding it are useful in the  
CC treatment of osteoarthritis, where they may be used for lubricating  
CC mammalian joints, such as articulating joints of humans, dogs or horses.  
CC The tribonectin, when formulated as a membrane, foam, gel or fibre, is  
CC useful for inhibiting adhesion between two surfaces such as the injured  
CC tissues of a mammal, where the injury is caused by a surgical insertion  
CC of trauma, or an artificial device e.g., an orthopaedic implant. In  
CC particular, one of the surfaces is pericardial tissue. DNA encoding a  
CC tribonectin may be used in gene therapy. The present sequence represents  
CC a human MSF gene reverse transcription-PCR (RT-PCR) primer used in an  
CC exemplification of the invention.  
XX  
XX Sequence 28 BP; 5 A; 7 C; 6 G; 10 T; 0 other;  
SQ  
Query Match 100.0%; Score 10; DB 22; Length 28;  
Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
Db |||||  
12 CTTCTCTTTT 21  
RESULT 97  
AAF05757/c  
ID AAF05757 standard; RNA; 29 BP.  
AC AAF05757;  
XX  
XX 16-FEB-2001 (first entry)  
DT  
XX Hammerhead ribozyme #2976.  
DE  
XX Ribozyme; erythropoietin; granulocyte colony stimulating factor;  
KW interferon alpha; ss.  
KW Homo sapiens.  
OS  
XX WO200061729-A2.  
PN  
XX 19-OCT-2000.  
PD  
XX 11-APR-2000; 2000WO-US09721.  
PF  
XX 12-APR-1999; 99US-0129390.  
XX

PA (RIBO-) RIBOZYME PHARM INC.  
XX Blatt L, Zwick M, Pavco P, McSwiggen J;  
PI WPI; 2000-647423/62.  
XX  
XX Enzymatic and antisense nucleic acid inhibition of repressor genes,  
PT useful for producing e.g. granulocyte colony stimulating factor  
PT protein, interferon alpha and erythropoietin -  
XX Claim 48; Page 124; 164pp; English.  
XX  
XX The present invention relates to enzymatic and antisense nucleic acid  
CC molecules that act as inhibitors of the expression of repressor genes  
CC encoding the TR2 Orphan receptor, EAR3/COUP-IF-1, the GATA  
CC transcription factor gene, IRF-2 and/or the CAAT Displacement  
CC Protein (CDP). Inhibition of the repressors removes prevents  
CC inhibition (and consequently increases expression of) genes involved in  
CC the production of erythropoietin, granulocyte colony stimulating factor  
CC protein and interferon alpha.  
XX  
XX Sequence 29 BP; 10 A; 4 C; 9 G; 5 U; 1 other;  
SQ  
Query Match 100.0%; Score 10; DB 21; Length 29;  
Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
Db |||||  
29 CTTCTCTTTT 20  
RESULT 98  
AAA40663  
ID AAA40663 standard; DNA; 29 BP.  
XX  
XX AAA40663;  
AC  
XX 15-AUG-2000 (first entry)  
DT  
XX Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:133.  
DE  
XX Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;  
KW therapy; screening; polymorphism; variant; detection; mutant;  
KW blood; mutation; insulin; glucose metabolism; fatty acid metabolism;  
KW catecholamine; malaria; infection; parasite; antiparasitic;  
KW antidiabetic; primer; ss.  
XX  
XX Homo sapiens.  
OS  
XX Synthetic.  
XX WO200019883-A2.  
PN  
XX 13-APR-2000.  
PD  
XX 07-OCT-1999; 99WO-US23418.  
PF  
XX 07-OCT-1998; 98US-0167750.  
PR  
XX 28-DEC-1998; 98US-0221222.  
PR  
XX 17-MAR-1999; 99US-0270542.  
PR  
XX (MEDI-) MEDICAL RES COUNCIL.  
PA (SCIO-) SCIOS INC.  
PA (AITM/) AITMAN T J.  
PA (SCOT/) SCOTT J.  
PA (STAN/) STANTON L W.  
XX  
XX Altman TJ, Scott J, Stanton LW;  
PI WPI; 2000-303596/26.  
XX  
XX Nucleic acids encoding mutant CD36 proteins useful for preventing,  
PT diagnosing and treating parasitic infections, especially malaria -  
PT



```

XX PS Claim 26; Page 90; 167pp; English.
XX CC The present invention describes isolated nucleic acid molecules (A)
CC encoding mutant CD36 proteins (B). Parasites such as Plasmodium
CC falciparum (the major cause of malaria) are unable to utilise the
CC mutated proteins to gain entry to, and infect cells. The mutant CD36
CC proteins do not function correctly preventing parasites utilising them
CC to infect cells. The nucleic acids may be used for the recombinant
CC production of mutant CD36 proteins according to standard methodologies.
CC They may be used in this way to prevent and treat parasitic infections
CC that utilise the CD36 protein to infect cells, such as P. falciparum,
CC the major cause of malaria. For example, the protein may be used to
CC identify modulators of CD36 expression and activity or a patient's CD36
CC DNA may be screened to determine whether there are any mutations present
CC that may confer resistance to parasitic infections. The proteins and
CC nucleic acids may also be used to prevent, diagnose and treat diseases
CC associated with defects in insulin action and/or glucose metabolism
CC and/or fatty acid metabolism and/or catecholamine action in subjects
CC possessing mutations in the CD36 genes. AAA40606 to AAA40759, and
CC AAB02515 to AAB02564, represent nucleotide and amino acid sequences
CC respectively which are used in the exemplification of the present
CC invention.
XX SQ Sequence 29 BP; 1 A; 5 C; 10 G; 13 T; 0 other;
Query Match 100.0%; Score 10; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17
RESULT 99
AAA40662
ID AAA40662 standard; DNA; 30 BP.
XX AC AAA40662;
XX DT 15-AUG-2000 (first entry)
XX DE Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:132.
XX KW Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;
XX KW therapy; screening; polymorphism; variant; detection; mutant;
XX KW blood; mutation; insulin; glucose metabolism; fatty acid metabolism;
XX KW catecholamine; malaria; infection; parasite; antiparasitic;
XX KW antidiabetic; primer; ss.
XX OS Homo sapiens.
XX OS Synthetic.
XX PN WO200019883-A2.
XX PD 13-APR-2000.
XX PF 07-OCT-1999; 99WO-US23418.
XX PR 07-OCT-1998; 98US-0167750.
XX PR 28-DEC-1998; 98US-0221222.
XX PR 17-MAR-1999; 99US-0270542.
XX PA (MEDI-) MEDICAL RES COUNCIL.
XX PA (SCIO-) SCIOS INC.
XX PA (AITM/) AITMAN T J.
XX PA (SCOT/) SCOTT J.
XX PA (STAN/) STANTON L W.
XX PI Aitman TJ, Scott J, Stanton LW;
XX WPI; 2000-303596/26.

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XX PT Nucleic acids encoding mutant CD36 proteins useful for preventing,
XX PT diagnosing and treating parasitic infections, especially malaria -
XX PS Claim 26; Page 90; 167pp; English.
XX CC The present invention describes isolated nucleic acid molecules (A)
XX encoding mutant CD36 proteins (B). Parasites such as Plasmodium
XX falciparum (the major cause of malaria) are unable to utilise the
XX mutated proteins to gain entry to, and infect cells. The mutant CD36
XX proteins do not function correctly preventing parasites utilising them
XX to infect cells. The nucleic acids may be used for the recombinant
XX production of mutant CD36 proteins according to standard methodologies.
XX They may be used in this way to prevent and treat parasitic infections
XX that utilise the CD36 protein to infect cells, such as P. falciparum,
XX the major cause of malaria. For example, the protein may be used to
XX identify modulators of CD36 expression and activity or a patient's CD36
XX DNA may be screened to determine whether there are any mutations present
XX that may confer resistance to parasitic infections. The proteins and
XX nucleic acids may also be used to prevent, diagnose and treat diseases
XX associated with defects in insulin action and/or glucose metabolism
XX and/or fatty acid metabolism and/or catecholamine action in subjects
XX possessing mutations in the CD36 genes. AAA40606 to AAA40759, and
XX AAB02515 to AAB02564, represent nucleotide and amino acid sequences
XX respectively which are used in the exemplification of the present
XX invention.
XX SQ Sequence 30 BP; 1 A; 5 C; 10 G; 14 T; 0 other;
Query Match 100.0%; Score 10; DB 21; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17
RESULT 100
AAA40664
ID AAA40664 standard; DNA; 30 BP.
XX AC AAA40664;
XX DT 15-AUG-2000 (first entry)
XX DE Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:134.
XX KW Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;
XX KW therapy; screening; polymorphism; variant; detection; mutant;
XX KW blood; mutation; insulin; glucose metabolism; fatty acid metabolism;
XX KW catecholamine; malaria; infection; parasite; antiparasitic;
XX KW antidiabetic; primer; ss.
XX OS Homo sapiens.
XX OS Synthetic.
XX PN WO200019883-A2.
XX PD 13-APR-2000.
XX PF 07-OCT-1999; 99WO-US23418.
XX PR 07-OCT-1998; 98US-0167750.
XX PR 28-DEC-1998; 98US-0221222.
XX PR 17-MAR-1999; 99US-0270542.
XX PA (MEDI-) MEDICAL RES COUNCIL.
XX PA (SCIO-) SCIOS INC.
XX PA (AITM/) AITMAN T J.
XX PA (SCOT/) SCOTT J.
XX PA (STAN/) STANTON L W.
XX

```

```

PI Aitman TJ, Scott J, Stanton LW;
XX
DR WPI; 2000-303596/26.
XX
PT Nucleic acids encoding mutant CD36 proteins useful for preventing,
PT diagnosing and treating parasitic infections, especially malaria -
XX
PS Claim 26; Page 90; 167pp; English.
XX
XX The present invention describes isolated nucleic acid molecules (A)
CC encoding mutant CD36 proteins (B). Parasites such as Plasmodium
CC falciparum (the major cause of malaria) are unable to utilise the
CC mutated proteins to gain entry to, and infect cells. The mutant CD36
CC proteins do not function correctly preventing parasites utilising them
CC to infect cells. The nucleic acids may be used for the recombinant
CC production of mutant CD36 proteins according to standard methodologies.
CC They may be used in this way to prevent and treat parasitic infections.
CC that utilise the CD36 protein to infect cells, such as P. falciparum,
CC the major cause of malaria. For example, the protein may be used to
CC identify modulators of CD36 expression and activity or a patient's CD36
CC DNA may be screened to determine whether there are any mutations present
CC that may confer resistance to parasitic infections. The proteins and
CC nucleic acids may also be used to prevent, diagnose and treat diseases
CC associated with defects in insulin action and/or glucose metabolism
CC and/or fatty acid metabolism and/or catecholamine action in subjects
CC possessing mutations in the CD36 genes. AAA40606 to AAA40759, and
CC AAB02515 to AAB02564, represent nucleotide and amino acid sequences
CC respectively which are used in the exemplification of the present
CC invention.
XX
SQ Sequence 30 BP; 1 A; 5 C; 10 G; 14 T; 0 other;

```

Query Match 100.0%; Score 10; DB 21; Length 30;  
Best Local Similarity 100.0%; Pred. No. 1.7e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

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Search completed: October 28, 2003, 17:17:58  
Job time : 287 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:14:10 ; Search time 2088 Seconds  
(without alignments)  
116.401 Million cell updates/sec

Title: US-09-335-032-71  
Perfect score: 10  
Sequence: 1 cttctctttt 10

Scoring table: OLIGO NUC  
Gapop 60.0 , Gapext 60.0

Searched: 22781392 seqs, 12152238056 residues

Word size : 0

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database :

- EST:\*  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hcc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hcc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vit:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rod:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrl:\*  
28: gb\_gss1:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	10	100.0	28	9	AI686998 tp81e01.x
3	10	100.0	28	29	TAI30812P
C 4	10	100.0	29	13	BQ590098

C 78	10	100.0	70	28	AZ918371	1006004B0	C 151	10	100.0	94	29	CC458005	CC458005_SALK_1147
C 79	10	100.0	71	14	D78209	D78209_EST	C 152	10	100.0	95	9	A1955060	A1955060_wg60c05.x
C 80	10	100.0	71	29	BZ770025	SALK_1429	C 153	10	100.0	95	9	AW713887	AW713887_h3jg02ne.f
C 81	10	100.0	72	9	AA184862	mu51b11.r	C 154	10	100.0	95	29	DR24C7S	DR24C7S_Danio rer
C 82	10	100.0	72	29	BX285944	Arabidops	C 155	10	100.0	96	13	RU863237	RU863237_S025F09.p
C 83	10	100.0	72	9	AA262253	zr70g04.r	C 156	10	100.0	96	28	AF179181	AF179181_AFI179181
C 84	10	100.0	73	9	AA97651	SWYD25CAU	C 157	10	100.0	96	29	BZ288800	BZ288800_SALK_0221
C 85	10	100.0	73	9	AW600116	SWL4CAK10	C 158	10	100.0	97	9	A1255137	A1255137_qv46d01.x
C 86	10	100.0	73	9	AW626514	SMOVL3CAN	C 159	10	100.0	97	9	A1360659	A1360659_qx59d01.x
C 87	10	100.0	73	9	AW626555	SMOVL3CAN	C 160	10	100.0	97	28	AZ461670	AZ461670_1M0267F11
C 88	10	100.0	73	9	AW651817	SWYD25CAU	C 161	10	100.0	97	29	BZ383278	BZ383278_SALK_1323
C 89	10	100.0	73	10	BG310475	SMOY3MCAU	C 162	10	100.0	97	29	CC029290	CC029290_3591_1.10
C 90	10	100.0	73	10	AW874933	SWYACAL04	C 163	10	100.0	97	29	AG229920	AG229920_Lotus_Tad
C 91	10	100.0	73	10	BE420470	SMOVL2CAS	C 164	10	100.0	98	10	BG621613	BG621613_601493758
C 92	10	100.0	73	10	BE420471	SMOVL2CAS	C 165	10	100.0	98	28	BH862099	BH862099_SALK_0887
C 93	10	100.0	73	10	BE420480	SMOVL2CAS	C 166	10	100.0	98	29	AL1770908	AL1770908_Arabidops
C 94	10	100.0	73	10	BEG38405	SMOVL2CAS	C 167	10	100.0	99	9	A1110549	A1110549_SMOVL3CAN
C 95	10	100.0	73	10	BF228818	SMOVL3CAN	C 168	10	100.0	99	9	AT006315	AT006315_AT006315
C 96	10	100.0	73	14	CB886667	SMOY3MCAU	C 169	10	100.0	99	28	AZ333242	AZ333242_1M0062D14
C 97	10	100.0	74	29	BZ660902	SALK_0243	C 170	10	100.0	99	29	CC459096	CC459096_SALK_1247
C 98	10	100.0	75	14	CA849383	K119503.y	C 171	10	100.0	100	9	AI197630	AI197630_ue45d06.r
C 99	10	100.0	75	29	TA77C02P	T. brucei	C 172	10	100.0	100	9	AW231456	AW231456_687061G10
C 100	10	100.0	76	9	AA615345	vo61d12.r	C 173	10	100.0	100	10	BG153513	BG153513_nag49f05.f
C 101	10	100.0	76	10	BF055674	7i68d03.y	C 174	10	100.0	100	10	BG153513	BG153513_nag49f05.f
C 102	10	100.0	76	28	AZ309821	1M0017C11	C 175	10	100.0	100	10	BE153449	BE153449_PM2-HT033
C 103	10	100.0	76	29	AL768539	Arabidops	C 176	10	100.0	100	13	BU097798	BU097798_946119D08
C 104	10	100.0	76	29	CNS02MOR	AL204372 Tetraodon	C 177	10	100.0	100	14	CB170638	CB170638_OVR602600
C 105	10	100.0	77	13	BQ456810	ke31c12.y	C 178	10	100.0	100	14	D82659	D82659_HUMHBC3277
C 106	10	100.0	77	14	CB099465	ks10h09.y	C 179	10	100.0	101	9	AL871534	AL871534_AL871534
C 107	10	100.0	77	14	CB277589	ks37g04.y	C 180	10	100.0	101	9	AA236344	AA236344_zr51f06.r
C 108	10	100.0	77	14	CB277809	ks39g02.y	C 181	10	100.0	101	10	BE500183	BE500183_WHE0980.F
C 109	10	100.0	77	29	AL758789	Arabidops	C 182	10	100.0	102	10	BF354509	BF354509_CM1-HT076
C 110	10	100.0	78	29	AL762583	Arabidops	C 183	10	100.0	102	12	BN090933	BN090933_1g17h06.x
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C 114	10	100.0	82	28	BH911702	SALK_0716	C 187	10	100.0	103	9	AW361423	AW361423_RC3-CT025
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C 117	10	100.0	83	14	F14596	S04D05 Por	C 190	10	100.0	103	12	BU327682	BU327682_RC3-BT033
C 118	10	100.0	83	14	F37812	HSPD06839 H	C 191	10	100.0	103	14	CA812476	CA812476_CA48LN061
C 119	10	100.0	83	29	AL941379	Arabidops	C 192	10	100.0	103	28	BH700682	BH700682_BOMMG89TF
C 120	10	100.0	84	14	CB001243	VVB002H02	C 193	10	100.0	103	28	BH818088	BH818088_BACP10-F
C 121	10	100.0	84	28	BH251470	SALK_0116	C 194	10	100.0	103	29	BZ314704	BZ314704_hz22e09.b
C 122	10	100.0	84	28	BH809934	SALK_0367	C 195	10	100.0	104	9	AI407102	AI407102_EST235390
C 123	10	100.0	84	29	CNS04INL	AL270426 Tetraodon	C 196	10	100.0	104	9	AA170839	AA170839_JTH131 HT
C 124	10	100.0	85	12	BI703760	ks18g12.y	C 197	10	100.0	104	9	AW062642	AW062642_RC0-CT009
C 125	10	100.0	85	13	BQ252968	sao04d04	C 198	10	100.0	104	10	BF747371	BF747371_RC3-BT033
C 126	10	100.0	85	14	U44279	ENU44279 As	C 199	10	100.0	104	10	BF833447	BF833447_PM1-HT092
C 127	10	100.0	85	28	AZ855489	2M0159L04	C 200	10	100.0	104	10	BF991035	BF991035_CM1-GN016
C 128	10	100.0	85	28	BH862052	SALK_0887	C 201	10	100.0	104	12	BI014338	BI014338_MR4-ET014
C 129	10	100.0	85	29	EX288399	Arabidops	C 202	10	100.0	104	12	BI044504	BI044504_PM3-OT020
C 130	10	100.0	86	9	AA067973	mm25a10.r	C 203	10	100.0	104	12	BI179001	BI179001_EST519946
C 131	10	100.0	86	9	AJ547920	AJ547920	C 204	10	100.0	104	12	BQ390565	BQ390565_BJ390565
C 132	10	100.0	86	10	BG673194	DRNBD05	C 205	10	100.0	104	12	BM403561	BM403561_zam4875.Z
C 133	10	100.0	86	29	BI175334	Danio rer	C 206	10	100.0	104	13	BQ358421	BQ358421_CM0-HT091
C 134	10	100.0	89	9	AV567495	AV567495	C 207	10	100.0	104	13	BQ864530	BQ864530_S04F12.F
C 135	10	100.0	89	28	BH251474	SALK_0116	C 208	10	100.0	104	29	CC027328	CC027328_3591_1.59
C 136	10	100.0	89	29	BZ292893	SALK_1285	C 209	10	100.0	105	9	AA068215	AA068215_mm48g02.r
C 137	10	100.0	90	13	BQ098870	ph23f03.y	C 210	10	100.0	105	9	AB036715	AB036715_AB036715
C 138	10	100.0	90	13	BQ582926	S015372-0	C 211	10	100.0	105	9	AA075027	AA075027_zm83h05.s
C 139	10	100.0	91	9	AA627648	ng51d08.s	C 212	10	100.0	105	9	AT006282	AT006282_AT006282
C 140	10	100.0	91	28	BH908214	SALK_0484	C 213	10	100.0	105	9	AA485070	AA485070_aa40b05.s
C 141	10	100.0	92	9	AA585277	PTH020 HT	C 214	10	100.0	105	13	BQ325343	BQ325343_CM1-CI000
C 142	10	100.0	92	14	C20817	HUMGS000486	C 215	10	100.0	105	14	CA798394	CA798394_Cac BL 72
C 143	10	100.0	92	28	BH215979	1006039C1	C 216	10	100.0	105	28	AZ797219	AZ797219_2M053015
C 144	10	100.0	92	29	CC179017	SALK_0571	C 217	10	100.0	106	9	AI230637	AI230637_EST227332
C 145	10	100.0	93	28	BH864683	SALK_0966	C 218	10	100.0	106	9	AU260000	AU260000_AU260000
C 146	10	100.0	93	28	BH864684	SALK_0967	C 219	10	100.0	106	9	AV902900	AV902900_AV902900
C 147	10	100.0	93	28	BH864686	SALK_0967	C 220	10	100.0	106	10	BE840762	BE840762_MR2-SN000
C 148	10	100.0	93	28	BH864689	SALK_0967	C 221	10	100.0	106	12	BI322353	BI322353_kx20a02.y
C 149	10	100.0	94	12	BI174502	OSTF046C1	C 222	10	100.0	106	14	CD026264	CD026264_NXSI_133
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224	10	100.0	107	12	BJ358573	BJ358573	297	10	100.0	115	10	BF475908	BF475908
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226	10	100.0	107	14	H23901	H23901	c 299	10	100.0	115	10	AW912033	AW912033
227	10	100.0	107	28	AZ565656	AZ565656	c 300	10	100.0	115	12	BM005397	BM005397
228	10	100.0	108	9	AW813619	AW813619	c 301	10	100.0	115	12	BM026664	BM026664
229	10	100.0	108	14	T83754	T83754	c 302	10	100.0	115	28	AQ357469	AQ357469
230	10	100.0	108	28	BH118434	BH118434	c 303	10	100.0	116	9	AI252189	AI252189
231	10	100.0	108	28	BH416075	BH416075	c 304	10	100.0	116	9	AW394059	AW394059
232	10	100.0	109	10	BE008587	BE008587	c 305	10	100.0	116	10	BG085551	BG085551
233	10	100.0	109	10	BE012104	BE012104	c 306	10	100.0	116	28	BH078074	BH078074
234	10	100.0	109	10	BE315587	BE315587	c 307	10	100.0	116	28	BZ080936	BZ080936
235	10	100.0	109	13	BQ591622	BQ591622	c 308	10	100.0	117	13	BH029480	BH029480
236	10	100.0	109	13	BQ405014	BQ405014	c 309	10	100.0	117	28	BI4642	BI4642
237	10	100.0	109	14	T91479	T91479	c 310	10	100.0	117	28	BH105487	BH105487
238	10	100.0	109	14	T91479	T91479	c 311	10	100.0	117	28	BZ377590	BZ377590
239	10	100.0	109	28	AQ080743	AQ080743	c 312	10	100.0	117	29	BZ764494	BZ764494
240	10	100.0	109	28	BH414927	BH414927	c 313	10	100.0	118	9	AA934372	AA934372
241	10	100.0	110	9	AA762509	AA762509	c 314	10	100.0	118	9	AW001605	AW001605
242	10	100.0	110	9	AV903734	AV903734	c 315	10	100.0	118	9	AW481540	AW481540
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244	10	100.0	110	14	CB006382	CB006382	c 317	10	100.0	118	10	AW845662	AW845662
245	10	100.0	110	28	BH347115	BH347115	c 318	10	100.0	118	10	BE998209	BE998209
246	10	100.0	110	28	BH637436	BH637436	c 319	10	100.0	118	12	BI400446	BI400446
247	10	100.0	110	28	BH809571	BH809571	c 320	10	100.0	118	14	CA881406	CA881406
248	10	100.0	110	28	AQ385208	AQ385208	c 321	10	100.0	118	14	D82788	D82788
249	10	100.0	110	29	CC365311	CC365311	c 322	10	100.0	118	14	T48514	T48514
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252	10	100.0	111	12	BM092640	BM092640	c 325	10	100.0	118	28	BH340928	BH340928
253	10	100.0	111	13	BM109777	BM109777	c 326	10	100.0	118	29	CNS0120E	CNS0120E
254	10	100.0	111	14	CB385906	CB385906	c 327	10	100.0	119	9	AV559784	AV559784
255	10	100.0	111	28	BZ115526	BZ115526	c 328	10	100.0	119	10	BF836144	BF836144
256	10	100.0	111	28	AQ482662	AQ482662	c 329	10	100.0	119	10	BG338662	BG338662
257	10	100.0	111	29	BZ290112	BZ290112	c 330	10	100.0	119	12	BM139880	BM139880
258	10	100.0	111	29	CC106912	CC106912	c 331	10	100.0	119	14	CA847224	CA847224
259	10	100.0	112	9	AA212505	AA212505	c 332	10	100.0	119	14	CB885142	CB885142
260	10	100.0	112	9	AW384207	AW384207	c 333	10	100.0	119	14	BM542039	BM542039
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262	10	100.0	112	9	BG604174	BG604174	c 335	10	100.0	119	29	AL754823	AL754823
263	10	100.0	112	12	BQ035562	BQ035562	c 336	10	100.0	120	9	AA082593	AA082593
264	10	100.0	112	14	R47206	R47206	c 337	10	100.0	120	9	AI998789	AI998789
265	10	100.0	112	28	AZ904406	AZ904406	c 338	10	100.0	120	9	AW393738	AW393738
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268	10	100.0	112	29	BZ734963	BZ734963	c 341	10	100.0	120	13	BU092901	BU092901
269	10	100.0	112	29	CC058942	CC058942	c 342	10	100.0	120	14	CD020233	CD020233
270	10	100.0	112	29	AL946882	AL946882	c 343	10	100.0	120	28	AZ736188	AZ736188
271	10	100.0	113	9	AV970176	AV970176	c 344	10	100.0	120	28	BH195988	BH195988
272	10	100.0	113	9	AW384162	AW384162	c 345	10	100.0	121	9	AA603854	AA603854
273	10	100.0	113	9	AW384195	AW384195	c 346	10	100.0	121	10	BE197817	BE197817
274	10	100.0	113	10	BF735099	BF735099	c 347	10	100.0	121	28	AZ086687	AZ086687
275	10	100.0	113	10	BE008812	BE008812	c 348	10	100.0	121	28	AZ879966	AZ879966
276	10	100.0	113	12	BJ364111	BJ364111	c 349	10	100.0	121	28	AZ911833	AZ911833
277	10	100.0	113	12	BJ369833	BJ369833	c 350	10	100.0	121	28	AZ918668	AZ918668
278	10	100.0	113	12	BJ409035	BJ409035	c 351	10	100.0	121	29	BZ958145	BZ958145
279	10	100.0	113	13	BQ114930	BQ114930	c 352	10	100.0	122	9	AA138752	AA138752
280	10	100.0	113	14	CA794594	CA794594	c 353	10	100.0	122	9	AW129644	AW129644
281	10	100.0	113	28	AZ564228	AZ564228	c 354	10	100.0	122	10	BF735541	BF735541
282	10	100.0	113	28	BH851383	BH851383	c 355	10	100.0	122	10	BF998919	BF998919
283	10	100.0	113	29	BZ358624	BZ358624	c 356	10	100.0	122	10	BG093701	BG093701
284	10	100.0	113	29	AL751707	AL751707	c 357	10	100.0	122	10	BG347903	BG347903
285	10	100.0	114	9	AL830940	AL830940	c 358	10	100.0	122	12	BG838320	BG838320
286	10	100.0	114	9	AI965770	AI965770	c 359	10	100.0	122	12	BI742650	BI742650
287	10	100.0	114	9	AW384183	AW384183	c 360	10	100.0	122	12	BJ363631	BJ363631
288	10	100.0	114	9	AW695613	AW695613	c 361	10	100.0	122	12	BU541349	BU541349
289	10	100.0	114	10	AW920975	AW920975	c 362	10	100.0	122	13	BU574463	BU574463
290	10	100.0	114	12	BJ538564	BJ538564	c 363	10	100.0	122	14	CB515109	CB515109
291	10	100.0	114	12	BM329382	BM329382	c 364	10	100.0	122	29	BX245707	BX245707
292	10	100.0	114	28	BH212127	BH212127	c 365	10	100.0	123	9	AA749641	AA749641
293	10	100.0	114	28	BH226094	BH226094	c 366	10	100.0	123	9	AU222440	AU222440
294	10	100.0	114	28	BH827449	BH827449	c 367	10	100.0	123	10	BF353080	BF353080
295	10	100.0	115	9	AI862639	AI862639	c 368	10	100.0	123	10	BF355020	BF355020
296	10	100.0	115	9	AV984486	AV984486	c 369	10	100.0	123	12	BI433964	BI433964



ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids ; eurosids II; Brassicales; Brassicaceae; Arabidopsiis.

REFERENCE  
AUTHORS 1 (bases 1 to 25)  
Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab , C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P., Zimmerman, J. and Ecker, J.R.

TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome

JOURNAL Unpublished

COMMENT Contact: Joseph R. Ecker  
Salk Institute Genomic Analysis Laboratory (SIGNAL)  
The Salk Institute for Biological Studies  
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
Tel: 858 453 4100 x1752  
Fax: 858 558 6379  
Email: ecker@salk.edu

TNNA. This is single pass sequence recovered from the left border of TNNA.

Class: TNNA tagged.  
Location/Qualifiers  
1. .25  
/organism="Arabidopsis thaliana"  
/mol\_type="genomic DNA"  
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/db\_xref="taxon:3702"  
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/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

BASE COUNT 16 a 1 c 7 g 1 t

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.5e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
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Db 22 CTTCTCTTTT 13

RESULT 2  
AI686998

LOCUS AI686998 28 bp mRNA linear EST 14-DEC-1999

DEFINITION tp81e01.x1 NCI CGAP Ut3 Homo sapiens cDNA clone IMAGE:2205720 3' similar to TR:054875 054875 MYOTONIC DYSTROPHY KINASE-RELATED CDC42-BINDING KINASE MRCK-BETA. ; contains element MER17 repetitive element ; mRNA sequence.

ACCESSION AI686998.1 GI:4898292

VERSION AI686998.1

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

AUTHORS 1 (bases 1 to 28)

TITLE NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>. National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)  
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: Greg Lennon, Ph.D.  
DNA Sequencing by: Washington University Genome Sequencing Center

Cloned through the I.M.A.G.E. Consortium/LNL at: [www-bio.llnl.gov/bbrp/image/image.html](http://www-bio.llnl.gov/bbrp/image/image.html)

Trace considered overall poor quality  
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Seq primer: -40UP from Gibco  
High quality sequence stop: 1.

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/lab\_host="DH10B"  
/clone\_lib="NCI CGAP Ut3"  
/note="Organ: uterus; Vector: pCMV-SPORT6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.45 kb. Life Technologies catalog #:  
11541-018"

BASE COUNT 4 a 12 c 0 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 28;  
Best Local Similarity 100.0%; Pred. No. 1.5e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||

Db 9 CTTCTCTTTT 18

RESULT 3  
TA130B12P

LOCUS TA130B12P 28 bp DNA linear GSS 13-DEC-2000

DEFINITION T. brucei sheared genomic DNA clone 130b12, forward sequence, genomic survey sequence.

ACCESSION AL464095

VERSION AL464095.1 GI:11834358

KEYWORDS GSS.

SOURCE Trypanosoma brucei

ORGANISM Trypanosoma brucei  
Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.

REFERENCE 1 (bases 1 to 28)

AUTHORS Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.E., Rajadream, M.A. and Barrell, B.G.

TITLE Direct Submission

JOURNAL Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: [barrell@sanger.ac.uk](mailto:barrell@sanger.ac.uk) and [nh@sanger.ac.uk](mailto:nh@sanger.ac.uk)

COMMENT Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTAT 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).  
Email: [nelsaved@tigr.org](mailto:nelsaved@tigr.org)  
Details of T. brucei sequencing at the Sanger Centre are available at [http://www.sanger.ac.uk/Projects/T\\_brucei/](http://www.sanger.ac.uk/Projects/T_brucei/).

FEATURES  
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Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 28

RESULT 4
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LOCUS
DEFINITION      BQ590098      29 bp      mRNA      linear      EST 06-DEC-2002
cDNA clone 024-019-019-T7 MP12-ADIS-024-storage root Beta vulgaris
ACCESSION      BQ590098
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Caryophyllales; Amaranthaceae; Beta.
REFERENCE
AUTHORS
Herwig,R., Schulz,B., Weisshaar,B., Hennig,S., Steinfath,M.,
Drungowski,M., Stahl,D., Wruck,W., Menze,A., O'Brien,J., Lehrach,H.
and Radelof,U.
TITLE
Construction of a 'unigene' cDNA clone set by oligonucleotide
fingerprinting allows access to 25 000 potential sugar beet genes
JOURNAL
Plant J. 32 (5), 845-857 (2002)
COMMENT
Contact: Weisshaar B
ADIS DNA core facility at MP12
Max-planck-Institute for Plant Breeding Research
Carl-von-Linne Weg 10, 50829 Koeln, Germany
Fax: 00492215062851
Email: weisshaar@mpiz-koeln.mpg.de
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Plate: 19 row: 0 column: 19
Seq primer: T7: GTAATACGACTCATTATAGGC.
FEATURES
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/db_xref="taxon:161934"
/clone="024-019-019"
/tissue_type="storage root"
/lab_host="EMDH10B"
/clone_lib="MP12-ADIS-024-storage root"
/notes="Vector: pCMVSPORT6; Site 1: Sali; Site 2: NotI;
cDNA library from sugar beet, library provided by KWS
Kleinwanzlebener Saatucht AG Einbeck, Germany, contact:
b.schulz@kws.de; cloning sites Sali-NotI, primer sites and
orientation:
SP6-Sali-CCAGCGTCG-5prime-cDNA-polyA-CC-NotI-T7; Note:
Sequencing granted in the context of the GABI-Beet project
, local PI: Dr. Katharina Schneider, coordinator: Prof.
Christian Jung; Sequence submission managed by
RZPD/GABI-Primary database: http://gabi.rzpd.de"
BASE COUNT      15 a      0 c      14 g      0 t
ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 1 CTTCTCTTTT 10

RESULT 5
BQ590098/c
LOCUS
DEFINITION      BQ590098      32 bp      mRNA      linear      EST 20-JUN-2001
cDNA clone 024-019-019-T7 MP12-ADIS-024-storage root Beta vulgaris
ACCESSION      BQ590098
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
NIH-MGC http://mgc.ncbi.nlm.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11059 row: e column: 23
High quality sequence stop: 32.
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Site 2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Library constructed by Life Technologies. Investigator
providing samples: Gilbert Smith, NIH"
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ORIGIN
Query Match      100.0%; Score 10; DB 12; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 20 CTTCTCTTTT 11

RESULT 6
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LOCUS
DEFINITION      AL941390      32 bp      DNA      linear      GSS 24-OCT-2002
Arabidopsis thaliana T-DNA flanking sequence GK-257B04-014921,
genomic survey sequence.
ACCESSION      AL941390
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
AUTHORS
Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
and Weisshaar,B.

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Db 18 CTTCTCTTTT 9

RESULT 5
BQ590098/c
LOCUS
DEFINITION      BQ590098      32 bp      mRNA      linear      EST 20-JUN-2001
cDNA clone 024-019-019-T7 MP12-ADIS-024-storage root Beta vulgaris
ACCESSION      BQ590098
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
NIH-MGC http://mgc.ncbi.nlm.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11059 row: e column: 23
High quality sequence stop: 32.
FEATURES
source
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/clone_lib="NCI CGAP Mam2"
/notes="Organ: mammary; Vector: pCMV-SPORT6; Site 1: Sali;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Library constructed by Life Technologies. Investigator
providing samples: Gilbert Smith, NIH"
BASE COUNT      19 a      5 c      8 g      0 t
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Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 20 CTTCTCTTTT 11

RESULT 6
AL941390
LOCUS
DEFINITION      AL941390      32 bp      DNA      linear      GSS 24-OCT-2002
Arabidopsis thaliana T-DNA flanking sequence GK-257B04-014921,
genomic survey sequence.
ACCESSION      AL941390
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
AUTHORS
Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
and Weisshaar,B.

```



**TITLE** A pipeline for automated high-throughput generation of ESTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines

**JOURNAL** Unpublished

**AUTHORS** Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weissshaar, B.

**TITLE** A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics

**JOURNAL** Unpublished

**REFERENCE** 3 (bases 1 to 32)

**AUTHORS** Rosso, M., Li, Y., Strizhov, N. and Weissshaar, B.

**JOURNAL** Direct Submission

**TITLE** Submitted (21-OCT-2002) Weissshaar B., Max-Planck-Institut fuer Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

**JOURNAL** This sequence is recovered from the left border of the T-DNA. It indicates an insertion close to or within gene At5g42900. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at: <http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

**FEATURES** Location/Qualifiers

1..32

    /organism="Arabidopsis thaliana"

    /mol\_type="genomic DNA"

    /strain="Columbia 0"

    /db\_xref="taxon:3702"

    /clone="GK-257B04-014921"

    /clone\_lib="Arabidopsis thaliana T-DNA insertion lines"

    /note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector pAC161. The lines contain one or more T-DNA from insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequences were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 0 a 6 c 0 g 26 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 32;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10

Db 10 CTCTCTTTT 19

RESULT 7

BZ763249

LOCUS SALK\_115724.25.30.n Arabidopsis thaliana T-DNA insertion lines

DEFINITION SALK\_115724.25.30.n Arabidopsis thaliana T-DNA insertion lines survey sequence.

ACCESSION BZ763249

VERSION BZ763249.1 GI:28935802

KEYWORDS GSS

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

REFERENCE 1 (bases 1 to 32)

AUTHORS Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prehn, L., Shinn, P., Zimmermann, J. and Ecker, J.R.

TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome

JOURNAL Unpublished

COMMENT Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies  
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
Tel: 858 453 4100 x1752  
Fax: 858 538 6379  
Email: ecker@salk.edu  
This is single pass sequence recovered from the left border of T-DNA.  
Class: T-DNA tagged.  
Location/Qualifiers

1..33

    /organism="Arabidopsis thaliana"

    /mol\_type="genomic DNA"

    /strain="Columbia 0"

    /db\_xref="taxon:3702"

    /clone="SALK\_115724.25.30.n"

    /clone\_lib="Arabidopsis thaliana T-DNA insertion lines"

    /note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more T-DNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

BASE COUNT 9 a 7 c 3 g 14 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 33;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10

Db 10 CTCTCTTTT 19

RESULT 8

AU256102/c

LOCUS AU256102 3'-directed mouse cDNA library Mus musculus cDNA clone

DEFINITION AU256102 3'-directed mouse cDNA library Mus musculus cDNA clone

ACCESSION AU256102

VERSION AU256102.1 GI:20319468

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 34)

AUTHORS Kato, K. and Matoba, R.

TITLE Generation of expressed sequence tags from mouse brain

JOURNAL Unpublished

COMMENT Contact: Kikuya Kato  
Graduate School of Biological Sciences  
Nara Institute of Science and Technology  
8916-5 Takayama, Ikoma, Nara 630-0101, Japan  
Tel: 81-743-72-5581  
Fax: 81-743-72-5589  
Email: kkatob@bs.nara.ac.jp, [love2.aist-nara.ac.jp/BED/index.html](mailto:love2.aist-nara.ac.jp/BED/index.html).

FEATURES Location/Qualifiers

1..34

    /organism="Mus musculus"

    /mol\_type="mRNA"

    /db\_xref="taxon:10090"

    /clone="BED0007468"

    /tissue="brain"

    /clone\_lib="3'-directed mouse cDNA library"

BASE COUNT 14 a 5 c 7 g 8 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 34;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 20 CTTCTCTTTT 11

RESULT 9  
 AZ429862  
 LOCUS  
 DEFINITION  
 ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT

AZ429862 37 bp DNA linear GSS 03-OCT-2000  
 1M0214105F Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
 clone UUGC1M0214105 F, genomic survey sequence.

AZ429862  
 AZ429862.1 GI:10553875  
 GSS  
 Mus musculus (house mouse)  
 Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 37)  
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,  
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.  
 and Wright, D., Weiss, R.  
 Mouse whole genome scaffolding with paired end reads from 10kb  
 plasmid inserts  
 Unpublished  
 Contact: Robert B. Weiss  
 University of Utah Genome Center  
 University of Utah  
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
 84112, USA  
 Tel: 801 585 5606  
 Fax: 801 585 7177  
 Email: ddunn@genetics.utah.edu  
 Insert Length: 10000 Std Error: 0.00  
 Plate: 0214 row: I column: 05  
 Seq primer: CGTTGTAAACGACGCCAGT  
 Class: plasmid ends  
 High quality sequence stop: 37.  
 Location/Qualifiers  
 1..37  
 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="UUGC1M0214105"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
 /clone\_lib="Mouse 10kb plasmid UUGC1M library"  
 /note="Vector: PWD42nv; Purified genomic DNA from M.  
 musculus C57BL/6J (male) was obtained from the Jackson  
 Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA  
 was hydrodynamically sheared by repeated passage through a  
 0.005 inch orifice at constant velocity. The sheared DNA  
 was blunt end-repaired with T4 DNA polymerase and T4  
 polynucleotide kinase. Adaptor oligonucleotides were  
 ligated to the blunt ends in high molar excess. The  
 adapted DNA was purified and size-selected for a 9.5 to  
 10.5 kb range using preparative agarose gel  
 electrophoresis. Vector DNA was prepared from a derivative  
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number  
 inducible derivative of plasmid R1. The vector was ligated  
 with adaptors complementary to the insert adaptors and  
 purified. The sheared, adapted mouse DNA was annealed to  
 adapted vector DNA, and transformed into  
 chemically-competent E. coli XL10-Gold (Stratagene) cells  
 and selected for ampicillin resistance."

FEATURES  
 source

BASE COUNT 8 a 9 c 6 g 14 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 28; Length 37;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
 |||||  
 Db 21 CTTCTCTTTT 30

RESULT 10  
 AZ514585  
 LOCUS  
 DEFINITION  
 ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT

AZ514585 37 bp DNA linear GSS 05-OCT-2000  
 1M0361B16F Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
 clone UUGC1M0361B16 F, genomic survey sequence.

AZ514585  
 AZ514585.1 GI:10695817  
 GSS  
 Mus musculus (house mouse)  
 Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 37)  
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,  
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.  
 and Wright, D., Weiss, R.  
 Mouse whole genome scaffolding with paired end reads from 10kb  
 plasmid inserts  
 Unpublished  
 Contact: Robert B. Weiss  
 University of Utah Genome Center  
 University of Utah  
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
 84112, USA  
 Tel: 801 585 5606  
 Fax: 801 585 7177  
 Email: ddunn@genetics.utah.edu  
 Insert Length: 10000 Std Error: 0.00  
 Plate: 0361 row: B column: 16  
 Seq primer: CGTTGTAAACGACGCCAGT  
 Class: plasmid ends  
 High quality sequence stop: 37.  
 Location/Qualifiers  
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 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="UUGC1M0361B16"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
 /clone\_lib="Mouse 10kb plasmid UUGC1M library"  
 /note="Vector: PWD42nv; Purified genomic DNA from M.  
 musculus C57BL/6J (male) was obtained from the Jackson  
 Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA  
 was hydrodynamically sheared by repeated passage through a  
 0.005 inch orifice at constant velocity. The sheared DNA  
 was blunt end-repaired with T4 DNA polymerase and T4  
 polynucleotide kinase. Adaptor oligonucleotides were  
 ligated to the blunt ends in high molar excess. The  
 adapted DNA was purified and size-selected for a 9.5 to  
 10.5 kb range using preparative agarose gel  
 electrophoresis. Vector DNA was prepared from a derivative  
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number  
 inducible derivative of plasmid R1. The vector was ligated  
 with adaptors complementary to the insert adaptors and  
 purified. The sheared, adapted mouse DNA was annealed to  
 adapted vector DNA, and transformed into  
 chemically-competent E. coli XL10-Gold (Stratagene) cells  
 and selected for ampicillin resistance."

FEATURES  
 source

BASE COUNT 3 a 8 c 2 g 24 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 28; Length 37;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 12 CTTCTCTTTT 21

RESULT 11  
 AZ613373  
 LOCUS  
 DEFINITION  
 1M0441M12R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
 clone UUGC1M0441M12 R, genomic survey sequence.

ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

1 (bases 1 to 39)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,  
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly  
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.  
 and Wright,D., Weiss,R.

TITLE  
 Mouse whole genome scaffolding with paired end reads from 10kb  
 plasmid inserts

JOURNAL  
 COMMENT  
 Contact: Robert B. Weiss  
 University of Utah Genome Center  
 University of Utah  
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
 84112, USA

FEATURES  
 source  
 1..39  
 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="UUGC1M0441M12"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, Tl-resistant, F-"  
 /clone\_lib="Mouse 10kb plasmid UUGC1M library"  
 /note="Vector: PWD42nv; Purified genomic DNA from M.  
 musculus C57BL/6J (male) was obtained from the Jackson  
 Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA  
 was hydrodynamically sheared by repeated passage through a  
 0.005 inch orifice at constant velocity. The sheared DNA  
 was blunt end-repaired with T4 DNA polymerase and T4  
 polynucleotide kinase. Adaptor oligonucleotides were  
 ligated to the blunt ends in high molar excess. The  
 adaptor DNA was purified and size-selected for a 9.5 to  
 10.5 kb range using preparative agarose gel  
 electrophoresis. Vector DNA was prepared from a derivative  
 of PWD42 (gi|4732114|gb|AF129072.1), a copy-number  
 inducible derivative of plasmid R1. The vector was ligated  
 with adaptors complementary to the insert adaptors and  
 purified. The sheared, adaptor mouse DNA was annealed to  
 adaptor vector DNA, and transformed into  
 chemically-competent E. coli XL10-Gold (Stratagene) cells  
 and selected for ampicillin resistance."

BASE COUNT  
 ORIGIN  
 8 a 8 c 2 g 21 t

Query Match 100.0%; Score 10; DB 28; Length 39;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 2 CTTCTCTTTT 11

RESULT 12  
 AA916182/c  
 LOCUS  
 DEFINITION  
 AA916182 40 bp mRNA linear EST 14-APR-1998  
 Q334B06.s1 NCI CGAP Br7 Homo sapiens cDNA clone IMAGE:1441715 3'  
 similar to TR:Q33575 Q33575 NADH DEHYDROGENASE SUBUNIT 4. ;, mRNA  
 sequence.

ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

1 (bases 1 to 40)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index

JOURNAL  
 COMMENT  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgapsb-remail.nih.gov  
 unknown library type

Trace considered overall poor quality  
 Seg primer: -40ml3 fwd. ET from Amersham  
 High quality sequence stop: 1.  
 Location/Qualifiers

FEATURES  
 source  
 1..40  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:1441715"  
 /lab\_host="DH10B"  
 /clone\_lib="NCI-CGAP Br7"  
 /note="Organ: breast; Vector: pCMV-SPORT4; Site 1: SalI;  
 Site 2: NotI; Cloned unidirectionally. Primer: Oligo dT.  
 Average insert size 1.2 kb. Life Technologies catalog  
 #:10985-018"

BASE COUNT  
 ORIGIN  
 26 a 0 c 12 g 2 t

Query Match 100.0%; Score 10; DB 9; Length 40;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 18 CTTCTCTTTT 9

RESULT 13  
 AZ375959/c  
 LOCUS  
 DEFINITION  
 AZ375959 41 bp DNA linear GSS 02-OCT-2000  
 1M0129A12R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
 clone UUGC1M0129A12 R, genomic survey sequence.

ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

1 (bases 1 to 41)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,  
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly  
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.  
 and Wright,D., Weiss,R.

BASE COUNT  
 ORIGIN  
 8 a 8 c 2 g 21 t

Query Match

**TITLE** Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts  
**JOURNAL** Unpublished  
**COMMENT** Contact: Robert B. Weiss  
 University of Utah  
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA  
 Tel: 801 585 5606  
 Fax: 801 585 7177  
 Email: ddunne@genetics.utah.edu  
 Insert Length: 10000 Std Error: 0.00  
 Plate: 0129 row: A column: 12  
 Seq primer: CACACAGGAACAGCTATGACC  
 Class: plasmid ends  
 High quality sequence stop: 41.

**FEATURES** source  
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 Location/Qualifiers  
 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="UUGCLM0129A12"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, Tl-resistant, F-"  
 /clone\_lib="Mouse 10kb plasmid UUGCLM library"  
 /note="Vector: pWb42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWb42 (G1.4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

**BASE COUNT** 22 a 4 c 7 g 8 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 28; Length 41;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**QY** 1 CTTCTCTTTT 10  
 |||||  
**Db** 25 CTTCTCTTTT 16

**RESULT 14**  
**EX120830/c**  
**LOCUS** BX120830 42 bp DNA linear GSS 13-MAR-2003  
**DEFINITION** Danio rerio genomic clone DKEY-61D1, genomic survey sequence.  
**ACCESSION** BX120830  
**VERSION** BX120830.1 GI:27951749  
**KEYWORDS** GSS.  
**SOURCE** Danio rerio (zebrafish)  
**ORGANISM** Danio rerio  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio.  
 1 (bases 1 to 42)  
 Humphray,S.J., Huckle,E. and Durham,J.L.  
 Direct Submission  
 Submitted (13-MAR-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished  
 This sequence was generated from the SP6 end of BAC 64G19. 64G19 is part of the Daniokey BAC library created by R. Plasterk and N.V. Keygene. Further details:  
 http://www.sanger.ac.uk/projects/D\_rerio/.

**FEATURES** source  
 1..42  
 Location/Qualifiers  
 /organism="Danio rerio"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:7955"  
 /clone="DKEY-61D1"  
 /tissue type="Testis"  
 /note="vector pIndigoBAC-536"  
 29 a 9 c 9 g 0 t

**BASE COUNT** 19 a 9 c 9 g 0 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 29; Length 42;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**QY** 1 CTTCTCTTTT 10  
 |||||  
**Db** 10 CTTCTCTTTT 1

**RESULT 16**  
**CC050354/c**  
**LOCUS** CC050354 45 bp DNA linear GSS 01-APR-2003  
**DEFINITION** OIS-536-6-lto6-G01 UniformMu MuTAIL Library Zea mays genomic clone OIS-536-6-lto6-G01, genomic survey sequence.

**COMMENT** Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished  
 This sequence was generated from the T7 end of BAC 61D1. 61D1 is part of the Daniokey BAC Library created by R. Plasterk and N.V. Keygene. Further details:  
 http://www.sanger.ac.uk/projects/D\_rerio/.

**FEATURES** source  
 1..42  
 Location/Qualifiers  
 /organism="Danio rerio"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:7955"  
 /clone="DKEY-61D1"  
 /tissue type="Testis"  
 /note="vector pIndigoBAC-536"  
 29 a 0 c 13 g 0 t

**BASE COUNT** 29 a 0 c 13 g 0 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 29; Length 42;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**QY** 1 CTTCTCTTTT 10  
 |||||  
**Db** 27 CTTCTCTTTT 18

**RESULT 15**  
**EX124544/c**  
**LOCUS** BX124544 42 bp DNA linear GSS 28-JAN-2003  
**DEFINITION** Danio rerio genomic clone DKEY-64G19, genomic survey sequence.  
**ACCESSION** BX124544  
**VERSION** BX124544.1 GI:27955482  
**KEYWORDS** GSS.  
**SOURCE** Danio rerio (zebrafish)  
**ORGANISM** Danio rerio  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio.  
 1 (bases 1 to 42)  
 Humphray,S.J., Huckle,E. and Durham,J.L.  
 Direct Submission  
 Submitted (27-JAN-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished  
 This sequence was generated from the SP6 end of BAC 64G19. 64G19 is part of the Daniokey BAC library created by R. Plasterk and N.V. Keygene. Further details:  
 http://www.sanger.ac.uk/projects/D\_rerio/.

**FEATURES** source  
 1..42  
 Location/Qualifiers  
 /organism="Danio rerio"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:7955"  
 /clone="DKEY-64G19"  
 /tissue type="Testis"  
 /note="vector pIndigoBAC-536"  
 19 a 9 c 9 g 0 t

**BASE COUNT** 19 a 9 c 9 g 0 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 29; Length 42;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**QY** 1 CTTCTCTTTT 10  
 |||||  
**Db** 10 CTTCTCTTTT 1

**RESULT 16**  
**CC050354/c**  
**LOCUS** CC050354 45 bp DNA linear GSS 01-APR-2003  
**DEFINITION** OIS-536-6-lto6-G01 UniformMu MuTAIL Library Zea mays genomic clone OIS-536-6-lto6-G01, genomic survey sequence.

```

ACCESSION      CC050354      GI:29465245
VERSION        CC050354.1
KEYWORDS       GSS.
SOURCE         Zea mays
ORGANISM       Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
Clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 45)
Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R.
Sequence tagged transposon insertions from the UniformMu maize
population
Unpublished
Contact: Donald R. McCarty
Plant Molecular and Cellular Biology Program
University of Florida
PO 110690 Gainesville, FL 32611-0690, USA
Tel: 352-392-1928 x322
Email: drmc@ufl.edu
Sequence flanking probable Mu insertion site in UniformMu line:
01S-536-6
Class: transposon insertion site.
Location/Qualifiers
1. .45
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="W22 (ACR, bz1-m9)"
/cultivar="UniformMu"
/db_xref="taxon:4577"
/clone="01S-536-6-1to6-G01"
/notes="Vector: TOPO-PCR4; DNA flanking Mu transposon
insertions in Mu inactive lines were extracted from the
UniformMu maize population by the thermo asymmetric
interlaced PCR (TAIL) protocol using primers specific for
the Mu terminal inverted repeat and a set of 16 arbitrary
primers. Amplicons were size enriched using Sepharose 400
spin columns and cloned into the TOPO PCR4 vector."
BASE COUNT    17 a 6 c 16 g 6 t
ORIGIN
Query Match    100.0%; Score 10; DB 29; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 20 CTTCTCTTTT 11

RESULT 17
LOCUS          BX120970
DEFINITION    Danio rerio genomic clone DKEY-68A12, genomic survey sequence.
ACCESSION     BX120970
VERSION       BX120970.1 GI:27951891
KEYWORDS      GSS.
SOURCE        Danio rerio (zebrafish)
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
1 (bases 1 to 45)
Humphray,S.J., Huckle,E. and Durham,J.L.
Submitted (27-JAN-2003)
Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humphray@sanger.ac.uk Unpublished
Direct Submission
Submitted (27-JAN-2003) The Sanger Institute, Wellcome Trust Genome
Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humphray@sanger.ac.uk Unpublished
This sequence was generated from the SP6 end of BAC 68A12. 68A12 is
part of the Daniokey BAC library created by R. Plasterk and N.V.
Keygene. Further details:
http://www.sanger.ac.uk/Projects/D_rerio/.
Location/Qualifiers

FEATURES
source
1. .45
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="DKEY-68A12"
/tissue_type="Testis"
/notes="Vector plndigoBAC-536"
0 t

BASE COUNT    28 a 2 c 15 g 0 t
ORIGIN
Query Match    100.0%; Score 10; DB 29; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 21 CTTCTCTTTT 12

RESULT 18
LOCUS          AA922976
DEFINITION    Ok77e06.s1 NCI CGAP GC4 Homo sapiens cDNA clone IMAGE1520002 3'
similar to TR:Q39614 Q39614 PROLINE-RICH PROTEIN. ;, mRNA sequence.
ACCESSION     AA922976
VERSION       AA922976.1 GI:3070285
KEYWORDS      EST.
SOURCE        Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 49)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. .49
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1520002"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/clone_lib="NCI-CGAP GC4"
/notes="Vector: pRT3D-Pac (Pharmacia) with a modified
polylinker; 1st strand cDNA was prepared from 3 pooled
germ cell tumors, and was then primed with a Not I -
oligo(dT) primer. Double-stranded cDNA was ligated to Eco
RI adaptors (Pharmacia), digested with Not I and cloned
into the Not I and Eco RI sites of the modified pRT3
vector. Library is normalized. Library was constructed by
Bento Soares and M. Fatima Bonaldo."
16 t

BASE COUNT    10 a 23 c 0 g 16 t
ORIGIN
Query Match    100.0%; Score 10; DB 9; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      1 CTTCTCTTTT 10
Db      1 CTTCTCTTTT 10

RESULT 19
A1564984
LOCUS   49 bp  mRNA  linear  EST 13-MAY-1999
DEFINITION
tq53b02.x1 NCI CGAP Utl1 Homo sapiens cDNA clone IMAGE:2212491 3'
similar to WP.F59E12.9 CELL1534 ;contains element MER22 repetitive
element ;, mRNA sequence.

ACCESSION
A1564984
VERSION 1
KEYWORDS
SOURCE  Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 49)
REFERENCE
AUTHORS  Alonso,J.M., Leisne,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
, C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.
, Zimmerman,J. and Ecker,J.R.
TITLE    A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL  Unpublished
COMMENT  Contact: Joseph R. Ecker
The Salk Institute Genomic Analysis Laboratory (SIGnAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: eckersalk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
1..49
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_130198.43.95.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 12 a 13 c 5 g 19 t
ORIGIN
Query Match 100.0%; Score 10; DB 29; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      1 CTTCTCTTTT 21

RESULT 21
A102691/c
LOCUS   50 bp  mRNA  linear  EST 30-AUG-2001
DEFINITION
HMA230056, mRNA sequence.

ACCESSION
A102691
VERSION 1
KEYWORDS
SOURCE  Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 50)
REFERENCE
AUTHORS  Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
, H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki
, Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE    Diverse transcriptional initiation revealed by fine, large-scale
mapping of mRNA start sites
JOURNAL  ENBO Rep. 2 (5), 388-393 (2001)
MEDLINE  21270072
PUBMED   11375929
COMMENT  Contact: Yutaka Suzuki
Department of Virology

FEATURES
source
1..49
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2212491"
/tissue_type="well-differentiated endometrial
adenocarcinoma, 7 pooled tumors"
/lab_host="DH10B"
/clone_lib="NCI CGAP Utl1"
/note="Organ: uterus; Vector: pCMV-SPORT6; Site 1: SalI;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.75 kb. Life Technologies catalog #:
11538-014"

BASE COUNT 0 a 26 c 0 g 23 t
ORIGIN
Query Match 100.0%; Score 10; DB 9; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      9 CTTCTCTTTT 18

RESULT 20
BZ357069
LOCUS   49 bp  DNA  linear  GSS 14-NOV-2002
DEFINITION
SALK_130198.43.95.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_130198.43.95.x, genomic
survey sequence.

ACCESSION
BZ357069

```

Institute of Medical Science, University of Tokyo  
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan  
Email: yasuku@ims.u-tokyo.ac.jp  
Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano  
S. Construction and characterization of a full length-enriched and  
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

# FEATURES

source

1. .50  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="HEM4230056"  
/clone\_lib="Sugano Homo sapiens cDNA library"

## BASE COUNT

ORIGIN

13 a 10 c 17 g 10 t

Query Match 100.0%; Score 10; DB 9; Length 50;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 28 CTTCTCTTTT 19

## RESULT 22

BZ770334

LOCUS

DEFINITION  
SALK\_143268.36.90.x Arabidopsis thaliana TDNA insertion lines  
Arabidopsis thaliana genomic clone SALK\_143268.36.90.x, genomic  
survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Arabisopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids  
; eurosids II; Brassicales; Brassicaceae; Arabidopsida.

REFERENCE

AUTHORS

Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadriab  
; Zimmermann, J. and Ecker, J.R.

TITLE

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished

COMMENT

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of

TDNA.

Class: TDNA tagged.

Location/Qualifiers

1. .51

/organism="Arabidopsis thaliana"

/mol\_type="genomic DNA"

/strain="Columbia 0"

/db\_xref="taxon:3702"

/clone="SALK\_143268.36.90.x"

/clone\_lib="Arabidopsis thaliana TDNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines

each of which contains one or more TDNA insertion

elements. The resultant fragment for each line was

directly sequenced to determine the genomic sequence at

the site of insertion. Details of the protocols used can

be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)

14 a 10 c 5 g 22 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 51;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 17 CTTCTCTTTT 26

## RESULT 23

AB082596

LOCUS

DEFINITION

Drosophila melanogaster DNA, clone:1(2)SH2 1592, genomic survey

sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Drosophila melanogaster (fruit fly)

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

AUTHORS

Oh, S., Kingsley, T., Shin, H., Zheng, Z., Chen, H. and Hou, S.

TITLE

Functional Genomics: A P element-mediated gene disruption in

Drosophila

Unpublished

JOURNAL

AUTHORS

Oh, S., Kingsley, T., Shin, H., Zheng, Z., Chen, H. and Hou, S.

TITLE

Direct Submission

JOURNAL

Submitted (26-MAR-2002) Suwan Oh, NCI-FCRDC, Lab. Of Immunobiology;  
1050 Boyles st., Frederick, Maryland 21702, USA  
(E-mail:ohsuwan@mail.ncifcrf.gov, Tel:1-301-846-7314,  
Fax:1-301-846-6145)

FEATURES

source

1. .51

/organism="Drosophila melanogaster"

/mol\_type="genomic DNA"

/db\_xref="taxon:7227"

/clone="1(2)SH2 1592"

BASE COUNT

ORIGIN

6 a 19 c 8 g 18 t

Query Match 100.0%; Score 10; DB 29; Length 51;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 28 CTTCTCTTTT 37

## RESULT 24

BQ667496/c

LOCUS

DEFINITION

Ancyllostoma caninum L3 serum scim pAMPI v1 Chiapelli McCarter

sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Ancyllostoma caninum (dog hookworm)

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;  
Ancylostomatidae; Ancylostomatinae; Ancylostominae; Ancylostoma.

REFERENCE

AUTHORS

McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,  
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,  
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Taagareishvili, R.,  
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe  
; Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,  
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and  
Wilson, R.

TITLE

The Washington Univ. Nematode EST Project, 1999

EST.

1 (bases 1 to 52)

McCart, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,  
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,  
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Taagareishvili, R.,  
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe  
; Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,  
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and  
Wilson, R.

TITLE

The Washington Univ. Nematode EST Project, 1999

BQ667496 52 bp mRNA linear EST 15-JUL-2002  
pb62c12.y1 Anc caninum L3 serum scim pAMPI v1 Chiapelli McCarter  
sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Ancylostoma caninum (dog hookworm)

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;  
Ancylostomatidae; Ancylostomatinae; Ancylostominae; Ancylostoma.  
REFERENCE  
1 (bases 1 to 52)

McCart, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,  
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,  
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Taagareishvili, R.,  
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe  
; Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,  
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and  
Wilson, R.

TITLE

The Washington Univ. Nematode EST Project, 1999

EST.

1 (bases 1 to 52)

McCart, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,  
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,  
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Taagareishvili, R.,  
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe  
; Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,  
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and  
Wilson, R.

TITLE

The Washington Univ. Nematode EST Project, 1999

JOURNAL  
COMMENT

Unpublished  
Contact: McCarter JP  
The Washington Univ. Nematode EST Project, 1999  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@wustl.edu  
The library was constructed by Brandi Chiappelli and Dr. James McCarter (bchiappell@wustl.edu & jmcarter@wustl.edu) at Washington University, St. Louis. DNA Sequencing by: Washington University Genome Sequencing Center St. Louis. Nematodes were provided by Dr. Prema Arasu of North Carolina State University. Putative full length read  
The vector to vector length is 53.

## FEATURES

source

1. 52  
Location/Qualifiers  
/organism="Nancylostoma caninum"  
/mol\_type="mRNA"  
/db\_xref="taxon:29170"  
/dev\_stage="serum stimulated L3"  
/lab\_host="DH10B"  
/clone\_lib="Anc caninum L3 serum stim pAMP1 v1 Chiappelli McCarter"  
/note="Vector: pAMP1 (Gibco); Site\_1: NotI; Site\_2: SalI; The library was constructed by Brandi Chiappelli and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna) PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of pAMP1. Nematodes were provided by Dr. Prema Arasu of North Carolina State University."

BASE COUNT 32 a 3 c 13 g 4 t  
ORIGIN

Query Match 100.0%; Score 10; DB 13; Length 52;

Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 44 CTTCTCTTTT 35  
|||||

## RESULT 25

AZ327146/c  
LOCUS

DEFINITION 52 bp DNA linear GSS 29-SEP-2000  
1M0050016F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0050016 F, genomic survey sequence.

ACCESSION AZ327146.1 GI:10385604

VERSION GSS.

KEYWORDS Mus musculus (house mouse)

SOURCE Mus musculus

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 52)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D.,Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

Unpublished

CONTACT: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0050 row: 0 column: 16

Seq primer: CGTTGTAAACGACGCGCCAGT

Class: plasmid ends

High quality sequence stop: 52.

Location/Qualifiers

source

1. 52  
/organism="Mus musculus"  
/mol\_type="genomic DNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUGC1M0050016"  
/sex="Male"  
/lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
/clone\_lib="Mouse 10kb plasmid UUGC1M library"  
/note="Vector: pWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (GI:4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 26 a 7 c 9 g 10 t  
ORIGIN

Query Match 100.0%; Score 10; DB 28; Length 52;

Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 32 CTTCTCTTTT 23  
|||||

## RESULT 26

BE569086/c

LOCUS

DEFINITION 55 bp mRNA linear EST 15-AUG-2000

601339390F2 NIH\_MGC\_53 Homo sapiens cDNA clone IMAGE:3681382 5', mRNA sequence.

ACCESSION BE569086

VERSION BE569086.1 GI:9812806

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 55)

NIH-MGC http://mgs.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished

CONTACT: Robert Strausberg, Ph.D.

Email: cgabbs@mail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: CLONTECH Laboratories, Inc.

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov

Plate: LLC363 row: d column: 23.

Location/Qualifiers

1. 52

## FEATURES

source



the site of insertion. Details of the protocols used can be found at <a href="http://signal.salk.edu/tdna_protocols.html">http://signal.salk.edu/tdna_protocols.html</a> "					
BASE COUNT	9 a	15 c	3 g	28 t	
ORIGIN					
Query Match	100.0%;	Score 10;	DB 28;	Length 55;	
Best Local Similarity	100.0%;	Pred. No. 1.6e+05;			
Matches	10;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	CTTCTCTTTT	10		
Db	42	CTTCTCTTTT	51		
RESULT 28					
BH864698/c					
LOCUS	BH864698 56 bp DNA linear GSS 05-AUG-2002				
DEFINITION	SALK_096733 Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_096733, genomic survey sequence.				
ACCESSION	BH864698				
VERSION	BH864698.1 GI:22100596				
KEYWORDS	GSS.				
SOURCE	Arabidopsis thaliana (thale cress)				
ORGANISM	Arabidopsis thaliana				
REFERENCE	1 (bases 1 to 56) Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrihab , C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P., Zimmerman,J. and Ecker,J.R. A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome				
JOURNAL	Unpublished				
COMMENT	Contact: Joseph R. Ecker Salk Institute Genomic Analysis Laboratory (SIGNAL) The Salk Institute for Biological Studies 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA Tel: 858 453 4100 x1752 Fax: 858 558 6379 Email: ecker@salk.edu This is single pass sequence recovered from the left border of TDNA.				
FEATURES	Class: TDNA tagged. Location/Qualifiers 1..56 /organism="Arabidopsis thaliana" /mol_type="genomic DNA" /strain="Columbia 0" /db_xref="taxon:3702" /clone="SALK_096733"				
BASE COUNT	19 a	5 c	12 g	20 t	
ORIGIN					
Query Match	100.0%;	Score 10;	DB 28;	Length 56;	
Best Local Similarity	100.0%;	Pred. No. 1.6e+05;			
Matches	10;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	CTTCTCTTTT	10		
Db	53	CTTCTCTTTT	44		
RESULT 29					
B1222307					
LOCUS	B1222307 57 bp mRNA linear EST 30-JUL-2002				

```

DEFINITION kx19c11.v3 Parastrongyloides trichosuri FL pAMP1 v1 Chiapelli
ACCESSION MCarter Parastrongyloides trichosuri cDNA 5', mRNA sequence.
VERSION BT322307
KEYWORDS BL322307.1 GI:15001493
SOURCE EST.
ORGANISM Parastrongyloides trichosuri
SOURCE Parastrongyloides trichosuri
ORGANISM Parastrongyloides trichosuri
REFERENCE 1 (bases 1 to 57)
AUTHORS McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,
Gibbons, M., Ritter, S., Bennett, J., Franklin, C., Tsagaris, V., R.,
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe
M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and
Wilson, R.
TITLE The Washington Univ. Nematode EST Project, 1999
JOURNAL Unpublished
COMMENT Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63103, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
The library was constructed by Brandi Chiapelli and Dr. James
McCarter (bchiapell@watson.wustl.edu & jmccarter@watson.wustl.edu) at
Washington University, St. Louis. DNA Sequencing by: Washington
University Genome Sequencing Center St. Louis.
Putative full length read
The vector to vector length is 58
Seq primer: -40RP from Gibco.
Location/Qualifiers
BASE COUNT 15 a 6 c 4 g 32 t
ORIGIN
/organism="Parastrongyloides trichosuri"
/mol_type="mRNA"
/db_xref="taxon:131310"
/dev_stage="Free Living"
/lab_host="DH10B"
/clone_lib="Parastrongyloides trichosuri FL pAMP1 v1
Chiapelli McCarter"
/notes="Vector: pAMP1 (Gibco); Site 1: NotI; Site 2: SalI;
The library was constructed by Brandi Chiapelli and Dr.
James McCarter at Washington University, St. Louis. The
cDNA was made by using Dynabead oligo-dT priming (Dyna).
PCR based library using a modified protocol from the
SMART PCR cDNA Synthesis Kit from Clontech. Directionally
cloned into the UDG sites of pAMP1. Nematodes were
provided by Dr. Warwick Grant of AgResearch, New Zealand
(warwick.grant@agresearch.co.nz)."
Query Match 100.0%; Score 10; DB 12; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTTT 10
|||||
Db 16 CTCTCTCTTTT 25

RESULT 30
CB917098 57 bp mRNA linear EST 25-APR-2003
LOCUS VVD118G10.371857 An expressed sequence tag database for abiotic
DEFINITION stressed berries of Vitis vinifera var. Chardonnay Vitis vinifera
cDNA clone VVD118G10 5, mRNA sequence.
ACCESSION CB917098
VERSION CB917098.1 GI:30131759
KEYWORDS EST.
SOURCE Vitis vinifera

DEFINITION tb92g09.x1 NCI CGAP Lu25 Homo sapiens cDNA clone IMAGE:2061856 3'
LOCUS AI343303 58 bp mRNA linear EST 08-APR-1999
DEFINITION similar to TR:Q33578 Q33578 KINETOPLAST CR5 ;, mRNA sequence.
ACCESSION AI343303
VERSION AI343303.1 GI:4080509
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 58)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
cDNA Library Preparation: David B. Krizman, Ph.D.
cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Trace considered overall poor quality

```

```

Seq primer: -40UP from Gibco
High quality sequence stop: 1.
FEATURES
    source
        1..58
        /organism="Homo sapiens"
        /mol_type="mRNA"
        /db_xref="taxon:9606"
        /clone="IMAGE:2061856"
        /tissue_type="bronchioalveolar carcinoma"
        /dev_stage="adult"
        /lab_host="DH10B"
        /clone_lib="NCI-GAP Lu25"
        /note="Organ: lung; Vector: pAMP1; mRNA made from lung
        carcinoma tissue, cDNA made by oligo-dr priming.
        Directionally cloned. Size-selected on agarose gel,
        average insert size 500 bp. Primary library,
        non-amplified. "
BASE COUNT      34 a      15 c      8 g      1 t
ORIGIN
    Query Match      100.0%; Score 10; DB 9; Length 58;
    Best Local Similarity 100.0%; Pred. No. 1.6e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
        |||||
        47 CTCTCTCTTT 38

Db

RESULT 32
LOCUS      BQ595228/c
DEFINITION      E012710-024-023-F23-SP6 MP1Z-ADIS-024-developing root Beta vulgaris
cDNA clone 024-023-F23 5-PRIME, mRNA sequence.
ACCESSION      BQ595228
VERSION      BQ595228.1 GI:26124811
KEYWORDS      EST.
SOURCE      Beta vulgaris
ORGANISM      Beta vulgaris
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Caryophyllales; Amaranthaceae; Beta.
REFERENCE      1 (bases 1 to 58)
AUTHORS      Herwig,R., Schulz,B., Weishaar,B., Hennig,S., Steinfath,M.,
Drungowski,M., Stahl,D., Wruck,W., Menze,A., O'Brien,J., Lehrach,H.
and Radelof,U.
TITLE      Construction of a 'unigenes' cDNA clone set by oligonucleotide
fingerprinting allows access to 25 000 potential sugar beet genes
Plant J. 32 (5), 845-857 (2002)
JOURNAL
COMMENT      Contact: Weishaar B
ADIS DNA core facility at MP1Z
Max-Planck-Institute for Plant Breeding Research
Carl-von-Linne Weg 10, 50829 Koeln, Germany
Fax: 00492215062851
Email: weissshaar@mpiz-koeln.mpg.de
Insert Length: 58 Std Error: 0.00
Plate: 23 row: F column: 23
Seq primer: SP6; CATACGATTAGTGACACTATAG.
FEATURES
    source
        1..58
        /organism="Beta vulgaris"
        /mol_type="mRNA"
        /cultivar="KWS2320 (double haploid, monogerm breeding line
        )"
        /db_xref="GABI:191621"
        /db_xref="taxon:161934"
        /clone="024-023-F23"
        /tissue_type="developing root"
        /lab_host="EMDH10B"
        /clone_lib="MP1Z-ADIS-024-developing root"
        /note="Vector: pCMVSPOR16; Site 1: SalI; Site 2: NotI;
        cDNA library from sugar beet, library provided by KWS
        Kleinwanzlebener Saatzzucht AG Binbeck, Germany, contact:

```

```

b.schulz@kws.de; cloning sites SalI-NotI, primer sites and
orientation:
SP6-Sali-CCACGGGTCCG-5prime-cDNA-polyA-CC-NotI-T7; Note:
Sequencing granted in the context of the GABI-Beet project
, local PI: Dr. Katharina Schneider, coordinator: Prof.
Christian Jung; Sequence submission managed by
RZPD/GABI-Primary database: http://gabi.rzpd.de"
BASE COUNT      32 a      8 c      17 g      1 t
ORIGIN
    Query Match      100.0%; Score 10; DB 13; Length 58;
    Best Local Similarity 100.0%; Pred. No. 1.6e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
        |||||
        54 CTCTCTCTTT 45

Db

RESULT 33
LOCUS      AU077187
DEFINITION      AU077187 Sugano cDNA library Homo sapiens cDNA clone Zrv61646
similar to 5'-end region of Mouse mRNA for proteasome Z subunit,
mRNA sequence.
ACCESSION      AU077187
VERSION      AU077187.1 GI:7439801
KEYWORDS      EST.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE      1 (bases 1 to 59)
AUTHORS      Suzuki,Y., Ishihara,D., Sasaki,M., Nakagawa,H., Hata,H., Tsunoda,T.,
Watanabe,M., Komatsu,T., Ota,T., Isogai,T., Suyama,A. and Sugano
,S.
TITLE      Statistical analysis of the 5' untranslated region of human mRNA
using 'Oligo-Capped' cDNA libraries
Genomics 64 (3), 286-297 (2000)
JOURNAL
MEDLINE      20221373
PUBMED      10756096
COMMENT      Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: ysuzuki@ims.u-tokyo.ac.jp
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
,S. Construction and characterization of a full length-enriched and
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997)
This clone was obtained from a '5'-end-enriched' cDNA library
constructed by 'Oligo-Capping' method. The coding region starts
from the 50 bp upstream to the 3'-end.
FEATURES
    source
        1..59
        /organism="Homo sapiens"
        /mol_type="mRNA"
        /db_xref="taxon:9606"
        /clone="Zrv61646"
        /clone_lib="Sugano cDNA library"
        /clone_size="19 g"
        /clone_start="12 c"
        /clone_end="19 t"
BASE COUNT      9 a      12 c      19 g      19 t
ORIGIN
    Query Match      100.0%; Score 10; DB 9; Length 59;
    Best Local Similarity 100.0%; Pred. No. 1.6e+05;
    Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
        |||||
        48 CTCTCTCTTT 57

Db

RESULT 34
AZ774339

```

**LOCUS**  
**DEFINITION** AZ774339 59 bp DNA linear GSS 16-FEB-2001  
 2M0003005R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
 clone UUGC2M0003005 R, genomic survey sequence.  
**ACCESSION** AZ774339  
**VERSION** GSS.  
**KEYWORDS** AZ774339.1 GI:12899665  
**SOURCE** Mus musculus (house mouse)  
**ORGANISM** Mus musculus  
**REFERENCE** Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
**AUTHORS** Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
 Islan, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,  
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.  
 and Wright, D., Weiss, R.  
**TITLE** Mouse whole genome scaffolding with paired end reads from 10kb  
 plasmid inserts  
**JOURNAL** Unpublished  
**COMMENT** Contact: Robert B. Weiss  
 University of Utah Genome Center  
 University of Utah  
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
 84112, USA  
 Tel: 801 585 5606  
 Fax: 801 585 7177  
 Email: dunn@genetics.utah.edu  
 Insert Length: 10000 Std Error: 0.00  
 Plate: 0003 row: 0 column: 05  
 Seq primer: CACACAGGAACAGCTATGACC  
 Class: plasmid ends  
 High quality sequence stop: 59.  
**FEATURES**  
 source  
 Location/Qualifiers  
 1..59  
 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="UUGC2M0003005"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, TI-resistant, F-"  
 /clone\_lib="Mouse 10kb plasmid UUGC1M library"  
 /notes="Vector: PWD42nv; Purified genomic DNA from M.  
 musculus C57BL/6J (male) was obtained from the Jackson  
 Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA  
 was hydrodynamically sheared by repeated passage through a  
 0.005 inch orifice at constant velocity. The sheared DNA  
 was blunt end-repaired with T4 DNA polymerase and T4  
 polynucleotide kinase. Adaptor oligonucleotides were  
 ligated to the blunt ends in high molar excess. The  
 adaptor DNA was purified and size-selected for a 9.5 to  
 10.5 Kb range using preparative agarose gel  
 electrophoresis. Vector DNA was prepared from a derivative  
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number  
 inducible derivative of plasmid R1. The vector was ligated  
 with adaptors complementary to the insert adaptors and  
 purified. The sheared, adaptor mouse DNA was annealed to  
 adaptor vector DNA, and transformed into  
 chemically-competent E. coli XL10-Gold (Stratagene) cells  
 and selected for ampicillin resistance."  
 12 a 7 c 12 g 28 t

**BASE COUNT** 12 a 7 c 12 g 28 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 28; Length 59;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 19 CTTCTCTTTT 29

RESULT 35

**LOCUS** BZ354535 59 bp DNA linear GSS 14-NOV-2002  
**DEFINITION** SALK\_125268.25.05.x Arabidopsis thaliana TDNA insertion lines  
 Arabidopsis thaliana genomic clone SALK\_125268.25.05.x, genomic  
 survey sequence.  
**ACCESSION** BZ354535  
**VERSION** BZ354535  
**KEYWORDS** GSS.  
**SOURCE** Arabidopsis thaliana (thale cress)  
**ORGANISM** Arabidopsis thaliana  
**REFERENCE** Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids  
 ; eurosids II; Brassicales; Brassicaceae; Arabidopsi.  
**AUTHORS** 1 (bases 1 to 59)  
 Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadriab  
 C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P.,  
 Zimmerman, J., and Ecker, J.R.  
**TITLE** A Sequence-Indexed Library of Insertion Mutations in the  
 Arabidopsis Genome  
**JOURNAL** Unpublished  
**COMMENT** Contact: Joseph R. Ecker  
 Salk Institute Genomic Analysis Laboratory (SIGnAL)  
 The Salk Institute for Biological Studies  
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
 Tel: 858 453 4100 x1752  
 Email: ecker@salk.edu  
 Fax: 858 558 6379  
 This is single pass sequence recovered from the left border of  
 TDNA.  
**Class:** TDNA tagged.

**FEATURES**  
 source  
 Location/Qualifiers  
 1..59  
 /organism="Arabidopsis thaliana"  
 /mol\_type="genomic DNA"  
 /strain="Columbia 0"  
 /db\_xref="taxon:3702"  
 /clone="SALK\_125268.25.05.x"  
 /clone\_lib="Arabidopsis thaliana TDNA insertion lines"  
 /notes="PCR was performed on Arabidopsis thaliana lines  
 each of which contains one or more TDNA insertion  
 elements. The resultant fragment for each line was  
 directly sequenced to determine the genomic sequence at  
 the site of insertion. Details of the protocols used can  
 be found at http://signal.salk.edu/tdna\_protocols.html"  
 7 a 16 c 14 g 22 t

**BASE COUNT** 7 a 16 c 14 g 22 t  
**ORIGIN**  
 Query Match 100.0%; Score 10; DB 29; Length 59;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Qy 1 CTTCTCTTTT 10  
 |||||  
 Db 1 CTTCTCTTTT 10

**RESULT 36**  
**AL762585**  
**LOCUS** AL762585 59 bp DNA linear GSS 19-JUN-2002  
**DEFINITION** Arabidopsis thaliana T-DNA flanking sequence GK-026B10-013759,  
 genomic survey sequence.  
**ACCESSION** AL762585  
**VERSION** AL762585.1 GI:21508841  
**KEYWORDS** GSS.  
**SOURCE** Arabidopsis thaliana (thale cress)  
**ORGANISM** Arabidopsis thaliana  
**REFERENCE** Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.  
**AUTHORS** 1  
 Strizhov, N., Li, Y., Rosso, M., Viehoveer, P., Dekker, K., Siedler, H.  
 and Weishaar, B.  
**TITLE** A pipeline for automated high-throughput generation of FSTs

(flanking sequence tags) from Arabidopsis thaliana T-DNA

transformed lines  
Unpublished

# JOURNAL REFERENCE

AUTHORS Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weishaar, B.  
TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics

# JOURNAL REFERENCE

AUTHORS Li, Y., Strizhov, N., Rosso, M. and Weishaar, B.  
TITLE Direct Submission  
JOURNAL Submitted (17-JUN-2002) Weishaar B., Max-Planck-Institut fuer Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, S0829, Germany  
COMMENT This sequence is recovered from the left border of the T-DNA. It indicates an insertion within the locus defined by clone F23B13. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at:  
<http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

# FEATURES

source

Location/Qualifiers

1..59

/organism="Arabidopsis thaliana"

/mol\_type="genomic DNA"

/strain="Columbia 0"

/db\_xref="taxon:3702"

/clone="GK-026B10-013759"

/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector PAC106. The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 9 a 11 c 12 g 27 t

# ORIGIN

Query Match

Best Local Similarity 100.0%; Score 10; DB 29; Length 59;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

33 CTTCTCTTTT 42

RESULT 37

AU258677

LOCUS

DEFINITION

AU258677 3'-directed mouse cDNA library Mus musculus cDNA clone BED0013467 3', mRNA sequence.

ACCESSION

AU258677

VERSION

KEYWORDS

SOURCE

ORGANISM

Mus musculus (house mouse)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Contact: Kikuya Kato  
Graduate School of Biological Sciences  
Nara Institute of Science and Technology  
8916-5 Takayama, Ikoma, Nara 630-0101, Japan  
Tel: 81-743-72-5581  
Fax: 81-743-72-5589  
Email: [kkato@bs.nara.ac.jp](mailto:kkato@bs.nara.ac.jp),  
URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

# FEATURES

source

Location/Qualifiers

1..63

/organism="Mus musculus"

/mol\_type="mRNA"

/db\_xref="taxon:10090"

/clone="BED0013467"

/tissue\_type="brain"

/clone\_lib="3'-directed mouse cDNA library"

16 a 11 c 11 g 24 t

1 others

Query Match

Best Local Similarity 100.0%; Score 10; DB 9; Length 63;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

15 CTTCTCTTTT 24

RESULT 38

AW874904/c

LOCUS

DEFINITION

AW874904

SWYACAL04A01SK Brugia malayi young adult cDNA (SAW99MLW-BmYA)

Brugia malayi cDNA clone SWYACAL04A01 5', mRNA sequence.

ACCESSION

AW874904.1

VERSION

AW874904.1

KEYWORDS

EST.

SOURCE

ORGANISM

Brugia malayi

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

Onchocercidae; Brugia.

REFERENCE

AUTHORS

WILLIAMS, S.A.

TITLE

Genes expressed in young adult of Brugia malayi

Unpublished

JOURNAL

COMMENT

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: [genome@smith.edu](mailto:genome@smith.edu)

Seq primer: pBluescript SK.

Location/Qualifiers

1..64

/organism="Brugia malayi"

/mol\_type="mRNA"

/db\_xref="taxon:6279"

/clone="SWYACAL04A01"

/dev\_stage="young adult"

/lab\_host="XLI-Blue MRF"

/clone\_lib="Brugia malayi young adult cDNA (SAW99MLW-BmYA)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Lymphatic filarial nematode parasite of humans. mRNA was prepared from young adult worms isolated from the peritoneal cavity of jirds and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 6.5 x 104 independent recombinants and the average insert size is approx. 800bp. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr. S.A. Williams, email: [genome@smith.edu](mailto:genome@smith.edu)."

25 a 10 c 18 g 11 t

Query Match

Best Local Similarity 100.0%; Score 10; DB 10; Length 64;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

```

Db          34 CTCTCTCTTT 25
|||||
RESULT 39
AW874934/c
LOCUS      64 bp      mRNA      linear      EST 22-MAY-2000
DEFINITION SWACAL04D04SK Brugia malayi young adult cDNA (SAW99MLM-BmYA)
            Brugia malayi cDNA clone SWACAL04D04 5', mRNA sequence.
ACCESSION  AW874934
VERSION     AW874934.1 GI:9012645
KEYWORDS   EST.
SOURCE     Brugia malayi
ORGANISM   Brugia malayi
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Brugia.
            1 (bases 1 to 64)
REFERENCE  1 Williams, S.A.
            Genes expressed in young adult of Brugia malayi
            JOURNAL  Unpublished
            COMMENT  Contact: Steven A. Williams
                    Smith College Department of Biological Sciences
                    Department of Biological Sciences, Clark Science Center, Smith
                    College, Northampton, MA, 01063, USA
                    Tel: 4135853826
                    Fax: 4135853786
                    Email: genome@smith.edu
                    Seq primer: pBluescript SK.
                    Location/Qualifiers
                    1..64
                    /organism="Brugia malayi"
                    /mol_type="mRNA"
                    /db_xref="taxon:6279"
                    /clone="SWACAL04D04"
                    /dev_stage="young adult"
                    /lab_host="XLI-Blue MRF"
                    /clone_lib="Brugia malayi young adult cDNA (SAW99MLM-BmYA)"
                    /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                    Xho I; Lymphatic filarial nematode parasite of humans. mRNA was
                    prepared from young adult worms isolated from the
                    peritoneal cavity of jirds and converted to
                    double-stranded cDNA using reverse transcriptase and
                    oligo(dT) followed by RNase H and DNA pol I. The library
                    has 6.5 x 104 independent recombinants and the average
                    insert size is approx. 800bp. The library was constructed
                    by Michelle Lizotte-Waniewski. The library is available
                    from Dr. S.A. Williams, email: genome@neal.smith.edu."
                    from Dr. S.A. Williams, email: genome@neal.smith.edu."
BASE COUNT  26 a 11 c 19 g 8 t
ORIGIN
Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 38 CTCTCTCTTT 29
|||||
RESULT 40
BE239248/c
LOCUS      64 bp      mRNA      linear      EST 11-JUL-2000
DEFINITION SWOV12CAS08D03SK Onchocerca volvulus L2 larvae cDNA (SAW98MLM-OvL2)
            Onchocerca volvulus cDNA clone SWOV12CAS08D03 5', mRNA sequence.
ACCESSION  BE239248
VERSION     BE239248.1 GI:9034224
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
ORGANISM   Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
            1 (bases 1 to 64)
REFERENCE  1 Williams, S.A.
            Genes expressed in L2 larvae of Onchocerca volvulus
            JOURNAL  Unpublished
            COMMENT  Contact: Steven A. Williams
                    Smith College Department of Biological Sciences
                    Department of Biological Sciences, Clark Science Center, Smith
                    College, Northampton, MA, 01063, USA
                    Tel: 4135853826
                    Fax: 4135853786
                    Email: genome@smith.edu
                    Seq primer: pBluescript SK.
                    Location/Qualifiers
                    1..64
                    /organism="Onchocerca volvulus"
                    /mol_type="mRNA"
                    /db_xref="taxon:6282"
                    /clone="SWOV12CAS08D03"
                    /dev_stage="L2"
                    /lab_host="XLI-Blue MRF"
                    /clone_lib="Onchocerca volvulus L2 larvae cDNA
                    (SAW98MLM-OvL2)"
                    /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                    Xho I; Filarial nematode parasite of humans. mRNA was
                    prepared from approximately 9,000 L2s isolated from
                    infected mosquitoes from Kumba, Cameroon and converted to
                    double-stranded cDNA using reverse transcriptase and
                    oligo(dT) followed by RNase H and DNA pol I. The library
                    has 7.3 x 10E4 independent recombinants and the average
                    insert size is approximately 1kb. The library was
                    constructed by Michelle Lizotte-Waniewski. The library is
                    available from Dr. S.A. Williams, email: genome@smith.edu."
                    available from Dr. S.A. Williams, email: genome@smith.edu."
BASE COUNT  24 a 9 c 19 g 12 t
ORIGIN
Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 13 CTCTCTCTTT 4
|||||
RESULT 41
BE239284/c
LOCUS      64 bp      mRNA      linear      EST 11-JUL-2000
DEFINITION SWOV12CAS08G03SK Onchocerca volvulus L2 larvae cDNA (SAW98MLM-OvL2)
            Onchocerca volvulus cDNA clone SWOV12CAS08G03 5', mRNA sequence.
ACCESSION  BE239284
VERSION     BE239284.1 GI:9034248
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
ORGANISM   Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
            1 (bases 1 to 64)
REFERENCE  1 Williams, S.A.
            Genes expressed in L2 larvae of Onchocerca volvulus
            JOURNAL  Unpublished
            COMMENT  Contact: Steven A. Williams
                    Smith College Department of Biological Sciences
                    Department of Biological Sciences, Clark Science Center, Smith
                    College, Northampton, MA, 01063, USA
                    Tel: 4135853826
                    Fax: 4135853786
                    Email: genome@smith.edu
                    Seq primer: pBluescript SK.
                    Location/Qualifiers
                    1..64
                    /organism="Onchocerca volvulus"

```

```

REFERENCE  1 (bases 1 to 64)
AUTHORS   Williams, S.A.
TITLE     Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL   Unpublished
COMMENT   Molecular Parasitology
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
            1..64
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            /mol_type="mRNA"
            /db_xref="taxon:6282"
            /clone="SWOV12CAS08D03"
            /dev_stage="L2"
            /lab_host="XLI-Blue MRF"
            /clone_lib="Onchocerca volvulus L2 larvae cDNA
            (SAW98MLM-OvL2)"
            /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
            Xho I; Filarial nematode parasite of humans. mRNA was
            prepared from approximately 9,000 L2s isolated from
            infected mosquitoes from Kumba, Cameroon and converted to
            double-stranded cDNA using reverse transcriptase and
            oligo(dT) followed by RNase H and DNA pol I. The library
            has 7.3 x 10E4 independent recombinants and the average
            insert size is approximately 1kb. The library was
            constructed by Michelle Lizotte-Waniewski. The library is
            available from Dr. S.A. Williams, email: genome@smith.edu."
            available from Dr. S.A. Williams, email: genome@smith.edu."
BASE COUNT  24 a 9 c 19 g 12 t
ORIGIN
Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 13 CTCTCTCTTT 4
|||||
RESULT 41
BE239284/c
LOCUS      64 bp      mRNA      linear      EST 11-JUL-2000
DEFINITION SWOV12CAS08G03SK Onchocerca volvulus L2 larvae cDNA (SAW98MLM-OvL2)
            Onchocerca volvulus cDNA clone SWOV12CAS08G03 5', mRNA sequence.
ACCESSION  BE239284
VERSION     BE239284.1 GI:9034248
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
ORGANISM   Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
            1 (bases 1 to 64)
REFERENCE  1 Williams, S.A.
            Genes expressed in L2 larvae of Onchocerca volvulus
            JOURNAL  Unpublished
            COMMENT  Contact: Steven A. Williams
                    Smith College Department of Biological Sciences
                    Department of Biological Sciences, Clark Science Center, Smith
                    College, Northampton, MA, 01063, USA
                    Tel: 4135853826
                    Fax: 4135853786
                    Email: genome@smith.edu
                    Seq primer: pBluescript SK.
                    Location/Qualifiers
                    1..64
                    /organism="Onchocerca volvulus"

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/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS08G03"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      25 a      9 c      19 g      11 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      14 CTTCTCTTTT 5

RESULT 42
BE239293/c
LOCUS      64 bp      mRNA      linear      EST 11-JUL-2000
DEFINITION Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ACCESSION BE239293
VERSION BE239293.1 GI:9034257
KEYWORDS EST.
SOURCE Onchocerca volvulus
ORGANISM Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. 64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS08G03"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
constructed by Michelle Lizotte-Waniewski. The library is

```

```

available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      29 a      11 c      15 g      9 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      47 CTTCTCTTTT 38

RESULT 43
BE636299/c
LOCUS      64 bp      mRNA      linear      EST 25-AUG-2000
DEFINITION Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ACCESSION BE636299
VERSION BE636299.1 GI:9919506
KEYWORDS EST.
SOURCE Onchocerca volvulus
ORGANISM Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. 64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS13H10"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      29 a      11 c      14 g      10 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      38 CTTCTCTTTT 29

RESULT 44
BE636308/c
LOCUS      64 bp      mRNA      linear      EST 25-AUG-2000

```



DEFINITION SWOVl2CASI4A09SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)  
Onchocerca volvulus cDNA clone SWOVl2CASI4A09 5', mRNA sequence.

ACCESSION BE636308  
VERSION BE636308.1 GI:9919515  
KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

COMMENT

Contact: Steven A. Williams

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVl2CASI4A09"

/dev\_stage="L2"

/lab\_host="Xl1-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10<sup>5</sup> independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

23 a 9 c 19 g 13 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 11 CTCTCTCTTT 2

RESULT 45

BE636320/c

LOCUS

DEFINITION SWOVl2CASI4C02SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)

Onchocerca volvulus cDNA clone SWOVl2CASI4C02 5', mRNA sequence.

ACCESSION BE636320

VERSION BE636320.1 GI:9919527

KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

COMMENT

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVl2CASI4C02"

/dev\_stage="L2"

/lab\_host="Xl1-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10<sup>5</sup> independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

26 a 8 c 19 g 11 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 16 CTCTCTCTTT 7

RESULT 46

BE636344/c

LOCUS

DEFINITION SWOVl2CASI4B08SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)

Onchocerca volvulus cDNA clone SWOVl2CASI4B08 5', mRNA sequence.

ACCESSION BE636344

VERSION BE636344.1 GI:9919551

KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

COMMENT

Contact: Steven A. Williams

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVl2CASI4B08"

/dev\_stage="L2"

/lab\_host="Xl1-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

FEATURES

source



/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 30 a 10 c 14 g 10 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 38 CTTCTCTTTT 29

RESULT 47  
BE636363/c  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

BE636363 64 bp mRNA linear EST 25-AUG-2000  
SWOVL2CASI4G07SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)  
Onchocerca volvulus cDNA clone SWOVL2CASI4G07 5', mRNA sequence.

BE636363  
EST  
Onchocerca volvulus

Onchocerca volvulus  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVL2CASI4G07"

/dev\_stage="L2"

/lab\_host="XLI-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10E4 independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 27 a 8 c 18 g 11 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 18 CTTCTCTTTT 9

RESULT 48

BE636369/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVL2CASI4H03"

/dev\_stage="L2"

/lab\_host="XLI-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10E4 independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 23 a 9 c 19 g 13 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 11 CTTCTCTTTT 2

RESULT 49

BE636392/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pluescript SK.

Location/Qualifiers

1..64

/organism="Onchocerca volvulus"

/mol\_type="mRNA"

/db\_xref="taxon:6282"

/clone="SWOVL2CASI4H03"

/dev\_stage="L2"

/lab\_host="XLI-Blue MRF"

/clone\_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10E4 independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 27 a 8 c 18 g 11 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.

REFERENCE  
1. (bases 1 to 64)  
AUTHORS  
Lizotte-Waniewski, M. and Williams, S.A.  
TITLE  
Genes expressed in adult male stage of *Onchocerca volvulus*  
JOURNAL  
Unpublished  
COMMENT  
Contact: Steven A. Williams  
Molecular Parasitology  
Smith College Department of Biological Sciences  
Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA  
Tel: 4135853826  
Fax: 4135853786  
Email: genome@smith.edu

Seq primer: pBluescript SK.

FEATURES  
source  
1. .64  
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/clone="SWOVAMCAQ09D08"  
/sex="male"  
/dev\_stage="adult"  
/lab\_host="XLI-Blue MRP"  
/clone\_lib="Onchocerca volvulus adult male cDNA  
(SAW98MLW-OvAM)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. Six adult  
male worms of *Onchocerca volvulus* were isolated from  
consenting patients and quick frozen. Adult male mRNA was  
converted to double-stranded cDNA using reverse  
transcriptase and oligo(dT) followed by RNase H and DNA  
pol I. The library has 2 x 10E5 independent recombinants  
and the average insert size is ~1100bp. The library was  
constructed by Michelle Lizotte-Waniewski with worms  
provided by Dr. Sara Lustigman. The library is available  
from Dr. Steven A. Williams, email: genome@smith.edu."

BASE COUNT 30 a 11 c 13 g 10 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
|||||  
Db 54 CTTCTCTTTT 45  
RESULT 50  
BE636430/c  
LOCUS  
DEFINITION  
SWOVLCAS1302SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)  
Onchocerca volvulus cDNA clone SWOVLCAS13C02 5', mRNA sequence.  
ACCESSION  
BE636430  
VERSION  
BE636430.1 GI:9919457  
KEYWORDS  
EST.  
SOURCE  
Onchocerca volvulus  
ORGANISM  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.  
REFERENCE  
1. (bases 1 to 64)  
AUTHORS  
Williams, S.A.  
TITLE  
Genes expressed in L2 larvae of *Onchocerca volvulus*  
JOURNAL  
Unpublished  
COMMENT  
Contact: Steven A. Williams  
Molecular Parasitology  
Smith College Department of Biological Sciences  
Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA  
Tel: 4135853826  
Fax: 4135853786  
Email: genome@smith.edu

Seq primer: pBluescript SK.

FEATURES  
source  
1. .64  
Location/Qualifiers  
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/clone="SWOVL2CAS13C02"  
/dev\_stage="L2"  
/lab\_host="XLI-Blue MRP"  
/clone\_lib="Onchocerca volvulus L2 larvae cDNA  
(SAW98MLW-OvL2)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and  
oligo(dT) followed by RNase H and DNA pol I. The library  
has 7.3 x 10E4 independent recombinants and the average  
insert size is approximately 1kb. The library was  
constructed by Michelle Lizotte-Waniewski. The library is  
available from Dr. S.A. Williams, email: genome@smith.edu."

BASE COUNT 28 a 9 c 15 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
|||||  
Db 33 CTTCTCTTTT 24  
RESULT 51  
BE636451/c  
LOCUS  
DEFINITION  
BE636451  
VERSION  
BE636451.1 GI:9919478  
KEYWORDS  
EST.  
SOURCE  
Onchocerca volvulus  
ORGANISM  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.  
REFERENCE  
1. (bases 1 to 64)  
AUTHORS  
Williams, S.A.  
TITLE  
Genes expressed in L2 larvae of *Onchocerca volvulus*  
JOURNAL  
Unpublished  
COMMENT  
Contact: Steven A. Williams  
Molecular Parasitology  
Smith College Department of Biological Sciences  
Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA  
Tel: 4135853826  
Fax: 4135853786  
Email: genome@smith.edu

Seq primer: pBluescript SK.

FEATURES  
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(SAW98MLW-OvL2)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and

Seq primer: pBluescript SK.

FEATURES  
source  
1. .64  
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(SAW98MLW-OvL2)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and

Seq primer: pBluescript SK.

FEATURES  
source  
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/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and

Seq primer: pBluescript SK.

FEATURES  
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Location/Qualifiers  
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/dev\_stage="L2"  
/lab\_host="XLI-Blue MRP"  
/clone\_lib="Onchocerca volvulus L2 larvae cDNA  
(SAW98MLW-OvL2)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and

Seq primer: pBluescript SK.

FEATURES  
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(SAW98MLW-OvL2)"  
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
Xho I; Filarial nematode parasite of humans. mRNA was  
prepared from approximately 9,000 L2s isolated from  
infected mosquitoes from Kumba, Cameroon and converted to  
double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 26 a 8 c 19 g 11 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
|||||  
Db 16 CTTCTCTTTT 7

RESULT 52  
BE636456/c  
LOCUS BE636456 64 bp mRNA linear EST 25-AUG-2000  
DEFINITION SMOVL2CAS13F038K Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)  
Onchocerca volvulus cDNA clone SMOVL2CAS13F03 5', mRNA sequence.  
ACCESSION BE636456  
VERSION BE636456.1 GI:9919483  
KEYWORDS EST.  
SOURCE Onchocerca volvulus  
ORGANISM Onchocerca volvulus  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.  
1 (bases 1 to 64)  
Williams, S.A.  
Genes expressed in L2 larvae of Onchocerca volvulus  
Unpublished  
Contact: Steven A. Williams  
Molecular Parasitology  
Smith College Department of Biological Sciences  
Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA  
Tel: 4135853826  
Fax: 4135853786  
Email: genome@smith.edu  
Seq primer: pBluescript SK.  
Location/Qualifiers  
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/notes="vector: Lambda Uni-ZAP XR; Site\_1: Eco RI; Site\_2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 29 a 13 c 13 g 13 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
|||||  
Db 41 CTTCTCTTTT 32

RESULT 53  
BE636464/c  
LOCUS BE636464 64 bp mRNA linear EST 25-AUG-2000  
DEFINITION SMOVL2CAS13G02SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)  
Onchocerca volvulus cDNA clone SMOVL2CAS13G02 5', mRNA sequence.  
ACCESSION BE636464  
VERSION BE636464.1 GI:9919491  
KEYWORDS EST.  
SOURCE Onchocerca volvulus  
ORGANISM Onchocerca volvulus  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.  
1 (bases 1 to 64)  
Williams, S.A.  
Genes expressed in L2 larvae of Onchocerca volvulus  
Unpublished  
Contact: Steven A. Williams  
Molecular Parasitology  
Smith College Department of Biological Sciences  
Department of Biological Sciences, Clark Science Center, Smith  
College, Northampton, MA, 01063, USA  
Tel: 4135853826  
Fax: 4135853786  
Email: genome@smith.edu  
Seq primer: pBluescript SK.  
Location/Qualifiers  
FEATURES  
source  
1..64  
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/clone="SMOVL2CAS13G02"  
/dev\_stage="L2"  
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/notes="vector: Lambda Uni-ZAP XR; Site\_1: Eco RI; Site\_2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 28 a 10 c 15 g 11 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 10; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
|||||  
Db 35 CTTCTCTTTT 26

RESULT 54  
BE636472/c  
LOCUS BE636472 64 bp mRNA linear EST 25-AUG-2000  
DEFINITION SMOVL2CAS13H01SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)  
Onchocerca volvulus cDNA clone SMOVL2CAS13H01 5', mRNA sequence.  
ACCESSION BE636472  
VERSION BE636472.1 GI:9919499  
KEYWORDS EST.  
SOURCE Onchocerca volvulus  
ORGANISM Onchocerca volvulus  
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
Onchocercidae; Onchocerca.  
1 (bases 1 to 64)  
Williams, S.A.  
Genes expressed in L2 larvae of Onchocerca volvulus

```

JOURNAL
COMMENT
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOV2L2CAS13H01"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr. S.A. Williams, email: genome@smith.edu."

BASE COUNT      26 a      7 c      19 g      12 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      21 CTTCTCTTTT 12

RESULT 55
BF118488/c
LOCUS
DEFINITION
SWOV13CAN69E12SK Onchocerca volvulus infective larva cDNA
(SAW94WL-OvL3) Onchocerca volvulus cDNA clone SWOV13CAN69E12 5',
mRNA sequence.
ACCESSION
BF118488
VERSION
BF118488.1 GI:10992964
KEYWORDS
EST.
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
REFERENCE
AUTHORS
Williams, S.A., Ju, W., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
TITLE
Genes expressed in infective third stage larvae of Onchocerca
JOURNAL
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"

JOURNAL
COMMENT
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"

JOURNAL
COMMENT
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
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/mol_type="mRNA"

/strain="Sierra Leone"
/db_xref="taxon:6282"
/clone="SWOV13CAN69E12"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus infective larva cDNA
(SAW94WL-OvL3)"
/note="Vector: Lambda UniZAP XR; Site 1: Eco RI; Site 2:
Xho I; Cutaneous filarial nematode parasite of humans.
mRNA was prepared from third stage infective larvae of
Onchocerca volvulus isolated from mosquitoes 10 days after
infection and converted to double stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNAPol I. The library had 1.8 x 10E5 independent
recombinants and average insert size was 900 base pairs.
The library was constructed by Wenhong Lu. The library is
available from Dr. S.A. Williams, email genome@smith.edu."

BASE COUNT      27 a      8 c      18 g      11 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      18 CTTCTCTTTT 9

RESULT 56
BI097426/c
LOCUS
DEFINITION
BI097426      64 bp      mRNA      linear      EST 25-JUN-2001
SWOV3MCAM63F09SK Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-OvML3) Onchocerca volvulus cDNA clone SWOV3MCAM63F09 5',
mRNA sequence.
ACCESSION
BI097426
VERSION
BI097426.1 GI:14549083
KEYWORDS
EST.
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
REFERENCE
AUTHORS
Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
TITLE
Genes expressed in molting L3 larvae of Onchocerca
JOURNAL
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
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/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-OvML3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3

```

in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10<sup>6</sup> independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)." 10 t

BASE COUNT 30 a 11 c 13 g 10 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 |||||  
 Db 40 CTCTCTCTTT 31

RESULT 57  
 BI142401/c  
 LOCUS  
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 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62B04 5',  
 mRNA sequence.

ACCESSION BI142401.1 GI:14624111  
 VERSION  
 KEYWORDS  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 1 (bases 1 to 64)  
 Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 Genes expressed in molting L3 larvae of Onchocerca volvulus  
 Unpublished  
 Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.  
 Location/Qualifiers

FEATURES  
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 /clone="SWOV3MCAM62B04"  
 /dev\_stage="molting L3"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA (SL96MLW-Ovml3)"  
 /note="vector: Lambda Uni-Zap XR; Site\_1: Eco RI; Site\_2: Xho I; Filarial nematode parasite of humans. Third-stage larvae, L3, were isolated from infected black flies in Cameroon (forest strain). The L3 were cultured in 20% FCS in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in culture. L3 of O. volvulus molt to fourth-stage larvae by day 5 in culture. mRNA was isolated from approximately 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3 in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10<sup>6</sup> independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams."

The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)." 12 t

BASE COUNT 27 a 10 c 15 g 12 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCTCTCTTT 10  
 |||||  
 Db 35 CTCTCTCTTT 26

RESULT 58  
 BI142408/c  
 LOCUS  
 DEFINITION SWOV3MCAM62B12SK Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62B12 5',  
 mRNA sequence.

ACCESSION BI142408.1 GI:14624118  
 VERSION  
 KEYWORDS  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 1 (bases 1 to 64)  
 Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 Genes expressed in molting L3 larvae of Onchocerca volvulus  
 Unpublished  
 Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.  
 Location/Qualifiers

FEATURES  
 source  
 1..64  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Kumba, Cameroons"  
 /db\_xref="taxon:6282"  
 /clone="SWOV3MCAM62B12"  
 /dev\_stage="molting L3"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA (SL96MLW-Ovml3)"  
 /note="vector: Lambda Uni-Zap XR; Site\_1: Eco RI; Site\_2: Xho I; Filarial nematode parasite of humans. Third-stage larvae, L3, were isolated from infected black flies in Cameroon (forest strain). The L3 were cultured in 20% FCS in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in culture. L3 of O. volvulus molt to fourth-stage larvae by day 5 in culture. mRNA was isolated from approximately 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3 in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10<sup>6</sup> independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)." 11 t

BASE COUNT 28 a 8 c 17 g 11 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 |||||  
 Db 30 CTCTCTCTTT 21

RESULT 59  
 B1142422/c  
 LOCUS  
 DEFINITION  
 SWOV3MCAM62D04SK Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62D04 5',  
 mRNA sequence.

ACCESSION B1142422 GI:14624132  
 VERSION B1142422.1  
 KEYWORDS Onchocerca volvulus  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 REFERENCE 1 (bases 1 to 64)  
 AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pbluescript SK.

FEATURES  
 source  
 1..64  
 Location/Qualifiers  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Kumba, Cameroons"  
 /db\_xref="taxon:6282"  
 /clone="SWOV3MCAM62D04"  
 /dev\_stage="molting L3"  
 /lab\_host="XL1-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
 Xho I; Filarial nematode parasite of humans. Third-stage  
 larvae, L3, were isolated from infected black flies in  
 Cameroon (forest strain). The L3 were cultured in 20% FCS  
 in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in  
 culture. L3 of O. volvulus molt to fourth-stage larvae by  
 day 5 in culture. mRNA was isolated from approximately  
 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3  
 in culture, and converted to double-stranded cDNA using  
 reverse transcriptase and oligo(dT) followed by RNase H  
 and DNA pol I. The library was constructed in the lambda  
 Uni-Zap XR vector and has 1 x 10E6 independent  
 recombinants and the average insert size is ~1200 bp. The  
 library was constructed by Sara Lustigman and Michelle  
 Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.  
 The library is available from Dr. Sara Lustigman (email:  
 slustig@nyc.org)."

BASE COUNT 28 a 9 c 15 g 12 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 |||||  
 Db 34 CTCTCTCTTT 25

RESULT 60  
 B1142452/c  
 LOCUS  
 DEFINITION

ACCESSION B1142452  
 VERSION B1142452.1  
 KEYWORDS Onchocerca volvulus  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 REFERENCE 1 (bases 1 to 64)  
 AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pbluescript SK.

FEATURES  
 source  
 1..64  
 Location/Qualifiers  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Kumba, Cameroons"  
 /db\_xref="taxon:6282"  
 /clone="SWOV3MCAM62F12"  
 /dev\_stage="molting L3"  
 /lab\_host="XL1-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
 Xho I; Filarial nematode parasite of humans. Third-stage  
 larvae, L3, were isolated from infected black flies in  
 Cameroon (forest strain). The L3 were cultured in 20% FCS  
 in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in  
 culture. L3 of O. volvulus molt to fourth-stage larvae by  
 day 5 in culture. mRNA was isolated from approximately  
 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3  
 in culture, and converted to double-stranded cDNA using  
 reverse transcriptase and oligo(dT) followed by RNase H  
 and DNA pol I. The library was constructed in the lambda  
 Uni-Zap XR vector and has 1 x 10E6 independent  
 recombinants and the average insert size is ~1200 bp. The  
 library was constructed by Sara Lustigman and Michelle  
 Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.  
 The library is available from Dr. Sara Lustigman (email:  
 slustig@nyc.org)."

BASE COUNT 27 a 10 c 15 g 12 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 |||||  
 Db 35 CTCTCTCTTT 26

RESULT 61  
 AZ821452  
 LOCUS

DEFINITION  
 2M0094BJ4F Mouse 10kb plasmid UUGCLM library Mus musculus genomic  
 clone UUGC2M0094B14 F, genomic survey sequence.  
 ACCESSION AZ821452  
 VERSION AZ821452.1 GI:12991360

B1142452 64 bp mRNA linear EST 05-JUL-2001  
 SWOV3MCAM62F12SK Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62F12 5',  
 mRNA sequence.

ACCESSION B1142452  
 VERSION B1142452.1  
 KEYWORDS Onchocerca volvulus  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 REFERENCE 1 (bases 1 to 64)  
 AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pbluescript SK.

FEATURES  
 source  
 1..64  
 Location/Qualifiers  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Kumba, Cameroons"  
 /db\_xref="taxon:6282"  
 /clone="SWOV3MCAM62F12"  
 /dev\_stage="molting L3"  
 /lab\_host="XL1-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-Ovml3)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:  
 Xho I; Filarial nematode parasite of humans. Third-stage  
 larvae, L3, were isolated from infected black flies in  
 Cameroon (forest strain). The L3 were cultured in 20% FCS  
 in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in  
 culture. L3 of O. volvulus molt to fourth-stage larvae by  
 day 5 in culture. mRNA was isolated from approximately  
 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3  
 in culture, and converted to double-stranded cDNA using  
 reverse transcriptase and oligo(dT) followed by RNase H  
 and DNA pol I. The library was constructed in the lambda  
 Uni-Zap XR vector and has 1 x 10E6 independent  
 recombinants and the average insert size is ~1200 bp. The  
 library was constructed by Sara Lustigman and Michelle  
 Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.  
 The library is available from Dr. Sara Lustigman (email:  
 slustig@nyc.org)."

BASE COUNT 27 a 10 c 15 g 12 t  
 ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10  
 |||||  
 Db 35 CTCTCTCTTT 26

RESULT 61  
 AZ821452  
 LOCUS  
 DEFINITION  
 2M0094BJ4F Mouse 10kb plasmid UUGCLM library Mus musculus genomic  
 clone UUGC2M0094B14 F, genomic survey sequence.  
 ACCESSION AZ821452  
 VERSION AZ821452.1 GI:12991360

KEYWORDS GSS.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 64)

REFERENCE  
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A. and Wright, D., Weiss, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL Unpublished

COMMENT Contact: Robert B. Weiss  
University of Utah Genome Center  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00  
Plate: 0094 row: B column: 14  
Seq primer: CTTTGTAAACGACGGCCACT  
Class: plasmid ends  
High quality sequence stop: 64.

FEATURES  
source  
1. .64  
Location/Qualifiers  
/organism="Mus musculus"  
/mol\_type="genomic DNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUGC2M0094B14"  
/sex="Male"  
/lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
/clone\_lib="Mouse 10kb plasmid UUGC1M library"  
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gi|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 12 c 12 g 26 t

ORIGIN  
Query Match 100.0%; Score 10; DB 28; Length 64;  
Best Local Similarity 100.0%; Pred. NO. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 16 CTTCTCTTTT 25

RESULT 62  
CB030041/c  
LOCUS  
DEFINITION  
65 bp mRNA linear EST 13-JAN-2003  
TgESTzyl8e02.y1 TgrH Tachyzoite Norm 7 cDNA Library Toxoplasma gondii cDNA clone TgESTzyl8e02.y1 5', mRNA sequence.

ACCESSION  
CB030041

CB030041.1 GI:2726413  
EST.  
Toxoplasma gondii  
Toxoplasma gondii  
Bukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida; Sarcocystidae; Toxoplasma.  
1 (bases 1 to 65)  
Tang, K., Cole, R., Fogarty, S., Sibley, L. D., Ajioke, J. A., White, M., Clifton, S., Pape, D., Martin, J., Wylie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y., Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagareishvili, R., Ronko, I., Kennedy, S., Maguire, L., Waterston, R. and Wilson, R.  
Toxoplasma EST Project  
Unpublished  
Contact: Clifton, S.  
Toxoplasma EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: toxo@watson.wustl.edu  
Contact David Sibley (toxost@borcim.wustl.edu) for further information relating to organism, libraries, or clone availability.  
Seq primer: -40RP from Gibco.

FEATURES  
source  
1. .85  
Location/Qualifiers  
/organism="Toxoplasma gondii"  
/mol\_type="mRNA"  
/strain="RH (Type I)"  
/db\_xref="taxon:5911"  
/clone="TgESTzyl8e02.y1"  
/dev\_stage="Tachyzoite"  
/lab\_host="DH10B (GeneHog, Invitrogen, Inc.)"  
/clone\_lib="TgrH Tachyzoite Norm 7 cDNA Library"  
/note="Vector: pBluscript SK-; Site 1: EcoRI; Site 2: XhoI; Toxoplasma RH strain tachyzoites were grown in human foreskin fibroblast cultures in vitro. The library was originally constructed by K. L. Wan, Cambridge University. cDNAs were synthesized from polyA RNAs by oligo d(T) priming and directionally cloned into the EcoRI to XhoI sites of the Lambda ZapII vector using the ZAP-cDNA synthesis kit (Stratagene). The primary cDNA library was mass excised as phagemid using ExAssist helper phage (Stratagene). Phagemid DNA was extracted by phenol-chloroform method, and hybridized against a pool of highly abundant genes which were derived from short-cycle PCR of the primary cDNA library. The normalized library was electroporated into DH10B (GeneHog, Invitrogen, Inc). WARNING: the library contains a small percentage of cDNAs derived from the human host cells."

BASE COUNT 31 a 7 c 20 g 7 t

ORIGIN  
Query Match 100.0%; Score 10; DB 14; Length 65;  
Best Local Similarity 100.0%; Pred. NO. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 58 CTTCTCTTTT 49

RESULT 63  
AZ514453/c  
LOCUS  
DEFINITION  
65 bp DNA linear GSS 05-OCT-2000  
IM0361N14F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0361N14 F, genomic survey sequence.

ACCESSION  
AZ514453  
VERSION  
GSS.  
KEYWORDS  
Mus musculus (house mouse)  
ORGANISM  
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;



ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids  
; eurosids II; Brassicales; Brassicaceae; Arabidopsids.  
REFERENCE 1 (bases 1 to 65)  
AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab,  
C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,  
Zimmerman,J. and Ecker,J.R.  
TITLE A Sequence-Indexed Library of Insertion Mutations in the  
Arabidopsis Genome  
JOURNAL Unpublished  
COMMENT Contact: Joseph R. Ecker  
Salk Institute Genomic Analysis Laboratory (SIGNAL)  
The Salk Institute for Biological Studies  
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
Tel: 858 453 4100 x1752  
Fax: 858 558 6379  
Email: ecker@salk.edu  
This is single pass sequence recovered from the left border of  
TDNA. This sequence lies within 300 bases of the 5' end of  
At5g15860.  
Class: TDNA tagged.  
Location/Qualifiers  
1. .65  
/organism="Mus musculus"  
/mol type="genomic DNA"  
/strain="C57BL/6J"  
/db xref="taxon:10090"  
/clones="UGGCM0361N14"  
/sex="Male"  
/lab host="E. Coli strain XL10-Gold, TI-resistant, F-"  
/clone lib="Mouse 10kb plasmid UGGCM library"  
/notes="Vector: PWD42nv; Purified genomic DNA from M.  
musculus C57BL/6J (male) was obtained from the Jackson  
Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA  
was hydrodynamically sheared by repeated passage through a  
0.005 inch orifice at constant velocity. The sheared DNA  
was blunt end-repaired with T4 DNA polymerase and T4  
polynucleotide kinase. Adaptor oligonucleotides were  
ligated to the blunt ends in high molar excess. The  
adaptored DNA was purified and size-selected for a 9.5 to  
10.5 kb range using preparative agarose gel  
electrophoresis. Vector DNA was prepared from a derivative  
of pWD42 (G114732114[gb|AF129072.1]), a copy-number  
inducible derivative of plasmid R1. The vector was ligated  
with adaptors complementary to the insert adaptors and  
purified. The sheared, adaptored mouse DNA was annealed to  
adaptored vector DNA, and transformed into  
chemically-competent E. coli XL10-Gold (Stratagene) cells  
and selected for ampicillin resistance."

## FEATURES

source

BASE COUNT 34 a 5 c 21 g 5 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 28; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 25 CTTCTCTTTT 16  
RESULT 64  
CC179177/c  
LOCUS CC179177 65 bp DNA linear GSS 02-MAY-2003  
DEFINITION SALK\_059169.41.80.x Arabidopsis thaliana TDNA insertion lines  
Arabidopsis thaliana genomic clone SALK\_059169.41.80.x, genomic  
survey sequence.  
ACCESSION CC179177  
VERSION CC179177.1 GI:30317728  
KEYWORDS GSS.  
SOURCE Arabidopsis thaliana (thale cress)

## REFERENCE

ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids  
; eurosids II; Brassicales; Brassicaceae; Arabidopsids.  
REFERENCE 1 (bases 1 to 65)  
AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab,  
C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,  
Zimmerman,J. and Ecker,J.R.  
TITLE A Sequence-Indexed Library of Insertion Mutations in the  
Arabidopsis Genome  
JOURNAL Unpublished  
COMMENT Contact: Joseph R. Ecker  
Salk Institute Genomic Analysis Laboratory (SIGNAL)  
The Salk Institute for Biological Studies  
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
Tel: 858 453 4100 x1752  
Fax: 858 558 6379  
Email: ecker@salk.edu  
This is single pass sequence recovered from the left border of  
TDNA. This sequence lies within 300 bases of the 5' end of  
At5g15860.  
Class: TDNA tagged.  
Location/Qualifiers  
1. .65  
/organism="Arabidopsis thaliana"  
/mol type="genomic DNA"  
/strain="Columbia 0"  
/db xref="taxon:3702"  
/clones="SALK\_059169.41.80.x"  
/clone lib="Arabidopsis thaliana TDNA insertion lines"  
/notes="PCR was performed on Arabidopsis thaliana lines  
each of which contains one or more TDNA insertion  
elements. The resultant fragment for each line was  
directly sequenced to determine the genomic sequence at  
the site of insertion. Details of the protocols used can  
be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

## FEATURES

source

BASE COUNT 33 a 7 c 14 g 11 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 29; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 41 CTTCTCTTTT 32  
RESULT 65  
AG216203  
LOCUS AG216203 65 bp DNA linear GSS 03-SEP-2002  
DEFINITION Drosophila melanogaster DNA, clone NP1611-5-1, flanking P[GawB]  
transposon insertion, genomic survey sequence.  
ACCESSION AG216203  
VERSION AG216203.1 GI:22763203  
KEYWORDS GSS.  
SOURCE Drosophila melanogaster (fruit fly)  
ORGANISM Drosophila melanogaster  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidea; Drosophilidae; Drosophila.  
REFERENCE 1  
AUTHORS Hayashi,S., Ito,K., Sado,Y., Taniguchi,M., Akimoto,A., Takeuchi,H.,  
Aigaki,T., Matsuzaki,F., Nakagoshi,H., Tanimura,T., Ueda,R.,  
Uemura,T., Yoshihara,M. and Goto,S.  
TITLE GEMDB, a database compiling expression patterns and molecular  
locations of a collection of Gal4 enhancer traps  
JOURNAL Genesis (2002) In press  
REFERENCE 2 (bases 1 to 65)  
AUTHORS Hayashi,S.  
TITLE Direct Submission  
JOURNAL Submitted (27-AUG-2002) Shigeo Hayashi, RIKEN Center for  
Developmental Biology, Laboratory for Morphogenetic Signaling;



Chuo-ku, Minatojima-minamimachi 2-2-3, Kobe, Hyogo 650-0047, Japan  
(E-mail: shayashi@cdb.riken.go.jp, Tel: 81-78-301-3184,  
Fax: 81-78-301-3183)

COMMENT This clone was isolated from genomic DNA flanking an insertion of the P element vector P[CaWB] of a *Drosophila* strain.

## FEATURES

source

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1. .65
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
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/chromosome="3"
/mapa="6288"
/clone="NP1611-5-1"
/note="flanking P[CaWB] transposon insertion"
BASE COUNT 6 a 34 c 4 g 20 t 1 others
ORIGIN
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Query Match 100.0%; Score 10; DB 29; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 41 CTTCTCTTTT 50

RESULT 66  
AL769814/c  
LOCUS  
DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-092H08-012001,  
genomic survey sequence.

ACCESSION AL769814  
VERSION AL769814.1 GI:21532016  
KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 Strizhov, N., Li, Y., Rosso, M., Viehoever, P., Dekker, K., Saedler, H.  
and Weisshaar, B.  
A pipeline for automated high-throughput generation of ESTs  
(flanking sequence tags) from Arabidopsis thaliana T-DNA  
transformed lines  
Unpublished

REFERENCE 2 Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weisshaar, B.  
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)  
for flanking sequence tag based reverse genetics  
Unpublished

JOURNAL Submitted (17-JUN-2002) Weisshaar B., Max-Planck-Institut fuer  
Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln 50829, Germany  
This sequence is recovered from the right border of the T-DNA. It  
indicates an insertion within the locus defined by clone MG46. The  
sequences are generated at the MPI for Plant Breeding Research in  
the context of the GABI-Kat project. GABI-Kat is part of the German  
Plant Genomics program designated 'GABI'. Information on line  
availability can be found at:  
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

## FEATURES

source

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1. .65
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-092H08-012001"
/note="PCR was performed on DNA from Arabidopsis thaliana
```

plants (T1) which were transformed with the T-DNA from vector pAC161. The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 32 a 9 c 11 g 13 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 29; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 39 CTTCTCTTTT 30

RESULT 67  
AL949773/c

LOCUS Arabidopsis thaliana T-DNA flanking sequence GK-323D01-015951,  
genomic survey sequence.

ACCESSION AL949773  
VERSION AL949773.1 GI:24406395  
KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 Strizhov, N., Li, Y., Rosso, M., Viehoever, P., Dekker, K., Saedler, H.  
and Weisshaar, B.  
A pipeline for automated high-throughput generation of ESTs  
(flanking sequence tags) from Arabidopsis thaliana T-DNA  
transformed lines  
Unpublished

REFERENCE 2 Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weisshaar, B.  
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)  
for flanking sequence tag based reverse genetics  
Unpublished

JOURNAL Submitted (21-OCT-2002) Weisshaar B., Max-Planck-Institut fuer  
Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln 50829, Germany  
This sequence is recovered from the left border of the T-DNA. It  
indicates an insertion within the locus defined by clone F9H3. The  
sequences are generated at the MPI for Plant Breeding Research in  
the context of the GABI-Kat project. GABI-Kat is part of the German  
Plant Genomics program designated 'GABI'. Information on line  
availability can be found at:  
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

REFERENCE 3 (bases 1 to 65)  
Strizhov, N., Li, Y., Rosso, M. and Weisshaar, B.  
Direct Submission

JOURNAL Submitted (21-OCT-2002) Weisshaar B., Max-Planck-Institut fuer  
Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln 50829, Germany  
This sequence is recovered from the left border of the T-DNA. It  
indicates an insertion within the locus defined by clone F9H3. The  
sequences are generated at the MPI for Plant Breeding Research in  
the context of the GABI-Kat project. GABI-Kat is part of the German  
Plant Genomics program designated 'GABI'. Information on line  
availability can be found at:  
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

## FEATURES

source

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1. .65
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-323D01-015951"
```

/clone lib="Arabidopsis thaliana T-DNA insertion lines"  
/notes="PCR was performed on DNA from Arabidopsis thaliana  
plants (T1) which were transformed with the T-DNA from  
vector pAC161. The lines contain one or more T-DNA  
insertions. The DNA fragment(s) resulting from the PCR  
were directly sequenced to determine the genomic sequence  
flanking the insertion. Sequences displaying significant  
similarity to the A. thaliana nuclear genome sequence were  
processed for submission. T-DNA derived sequences were

```

removed"
BASE COUNT      26 a      7 c      17 g      15 t
ORIGIN

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 12 CTTCTCTTTT 3

RESULT 68
TAL29A12P/c
LOCUS      TAL29A12P      65 bp      DNA      linear      GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 129a12, forward sequence,
            genomic survey sequence.
ACCESSION  AL463978
VERSION     AL463978.1 GI:11834241
KEYWORDS   GSS.
SOURCE     Trypanosoma brucei
ORGANISM   Trypanosoma brucei
            Eukaryota; Euzlenozoa; Kinetoplastida; Trypanosomatidae;
            Trypanosoma.
REFERENCE  1 (bases 1 to 65)
AUTHORS   Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
            Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
            Melville, S.E., Rajandream, M.A. and Barrell, B.G.
TITLE     Direct Submission
JOURNAL    Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
            project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
            Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
            nh@sanger.ac.uk
COMMENT    Constructed at the Institute for Genomic Research (TIGR),
            Rockville, MD. Genomic DNA isolated from a cloned population of
            Trypanosoma brucei (TREU27/4 GUTat 10.1) was mechanically sheared
            to give a tight size distribution (
            4 kb). The v + i method used for the library construction is
            described in detail in Smith, H. and Venter, J.C. (Making small
            insert libraries for whole genome shotgun sequencing projects. In
            Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
            Barrell, Oxford University Press, 1999).
            Email: nelsayed@tigr.org
            Details of T. brucei sequencing at the Sanger Centre are available
            at http://www.sanger.ac.uk/Projects/T_brucei/.
            Location/Qualifiers
                1..65
                /organism="Trypanosoma brucei"
                /mol_type="genomic DNA"
                /strain="TREU27"
                /db_xref="taxon:5691"
                /clone="129a12"

BASE COUNT      31 a      7 c      22 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 62 CTTCTCTTTT 53

RESULT 69
W85242/c
LOCUS      W85242      66 bp      mRNA      linear      EST 12-SEP-1996
DEFINITION m52h08.r1 Soares mouse embryo NBME13.5 14.5 Mus musculus cDNA
            clone IMAGE:408735 5' similar to gb:U13546 rnal Human HMG-17 gene
            for non-histone chromosomal protein (HUMAN); gb:U12944 Mouse mRNA
            for HMG-17 chromosomal protein (MOUSE); mRNA sequence.
W85242
ACCESSION

removed"
BASE COUNT      26 a      7 c      17 g      15 t
ORIGIN

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 12 CTTCTCTTTT 3

RESULT 68
TAL29A12P/c
LOCUS      TAL29A12P      65 bp      DNA      linear      GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 129a12, forward sequence,
            genomic survey sequence.
ACCESSION  AL463978
VERSION     AL463978.1 GI:11834241
KEYWORDS   GSS.
SOURCE     Trypanosoma brucei
ORGANISM   Trypanosoma brucei
            Eukaryota; Euzlenozoa; Kinetoplastida; Trypanosomatidae;
            Trypanosoma.
REFERENCE  1 (bases 1 to 65)
AUTHORS   Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
            Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
            Melville, S.E., Rajandream, M.A. and Barrell, B.G.
TITLE     Direct Submission
JOURNAL    Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
            project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
            Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
            nh@sanger.ac.uk
COMMENT    Constructed at the Institute for Genomic Research (TIGR),
            Rockville, MD. Genomic DNA isolated from a cloned population of
            Trypanosoma brucei (TREU27/4 GUTat 10.1) was mechanically sheared
            to give a tight size distribution (
            4 kb). The v + i method used for the library construction is
            described in detail in Smith, H. and Venter, J.C. (Making small
            insert libraries for whole genome shotgun sequencing projects. In
            Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
            Barrell, Oxford University Press, 1999).
            Email: nelsayed@tigr.org
            Details of T. brucei sequencing at the Sanger Centre are available
            at http://www.sanger.ac.uk/Projects/T_brucei/.
            Location/Qualifiers
                1..65
                /organism="Trypanosoma brucei"
                /mol_type="genomic DNA"
                /strain="TREU27"
                /db_xref="taxon:5691"
                /clone="129a12"

BASE COUNT      31 a      7 c      22 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 62 CTTCTCTTTT 53

RESULT 69
W85242/c
LOCUS      W85242      66 bp      mRNA      linear      EST 12-SEP-1996
DEFINITION m52h08.r1 Soares mouse embryo NBME13.5 14.5 Mus musculus cDNA
            clone IMAGE:408735 5' similar to gb:U13546 rnal Human HMG-17 gene
            for non-histone chromosomal protein (HUMAN); gb:U12944 Mouse mRNA
            for HMG-17 chromosomal protein (MOUSE); mRNA sequence.
W85242
ACCESSION

us-09-335-032-71.oli.rst
VERSION      W85242.1 GI:1397731
KEYWORDS     EST.
SOURCE       Mus musculus (house mouse)
ORGANISM     Mus musculus

REFERENCE
AUTHORS
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
Waterston, R.
TITLE        The WashU-HHMI Mouse EST Project
JOURNAL      Unpublished
COMMENT      Contact: Marra M/Mouse EST Project
            WashU-HHMI Mouse EST Project
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:252503
            Trace considered overall poor quality
            Seq primer: -28M13 rev2 from Amerisham
            High quality sequence stop: 1.
            Location/Qualifiers
                1..66
                /organism="Mus musculus"
                /mol_type="mRNA"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="IMAGE:408735"
                /sex="unknown"
                /tissue_type="embryo"
                /dev_stage="13.5-14.5dpc total fetus"
                /lab_host="DH10B"
                /clone_lib="Soares mouse embryo NBME13.5 14.5"
                /note="vector: pT73D-Pac (Pharmacia) with a modified
                polylinker; Site 1: Not 1; Site 2: Eco RI; 1st strand cDNA
                was primed with a Not I - oligo(gt) primer (5'
                TGTTCACATCTGAAGTGGGCGCGCGGAAATTTTTTTTTTTTTTTT
                T 3'), on equal amounts of mRNA from 2 13.5dpc and 2
                14.5dpc embryos [total RNA provided by Minoru Ko, Wayne
                State Univ., from 2 ]; double-stranded cDNA was ligated to
                Eco RI adaptors (Pharmacia), digested with Not I and
                cloned into the Not I and Eco RI sites of the modified
                pT73 vector. Library went through one round of
                normalization, and was constructed by Bento Soares and
                M. Patricia Bonaldo. "
                19 a      19 c      18 g      10 t

BASE COUNT
ORIGIN

Query Match      100.0%; Score 10; DB 14; Length 66;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 41 CTTCTCTTTT 32

RESULT 70
A2514401
LOCUS      A2514401
DEFINITION 1M0361X04F Mouse 10kb plasmid UUGCLM library Mus musculus genomic
            clone UUGC1M0361X04 F, genomic survey sequence.
ACCESSION  A2514401
VERSION     A2514401.1 GI:10695717
KEYWORDS   GSS
SOURCE     Mus musculus (house mouse)
ORGANISM   Mus musculus

```

Rukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 66)

**REFERENCE**  
Dunn D., Aoyagi A., Barber M., Beacorn T., Duval B., Hamil C., Islam H., Longacre S., Mahmoud M., Meenen E., Pedersen T., Reilly M., Rose M., Rose R., Stokes R., Tingey A., von Niedehausen A., and Wright D., Weiss R.

**AUTHORS**  
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

**TITLE**  
Unpublished

**JOURNAL**  
Contact: Robert B. Weiss

**COMMENT**  
University of Utah  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00  
Plate: 0361 row: K column: 04  
Seq primer: CGTGTAAACGACGGCCAGT  
Class: plasmid ends  
High quality sequence stop: 66.

**FEATURES**  
Location/Qualifiers  
1..66  
/organism="Mus musculus"  
/mol\_type="genomic DNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUGC1M0361K04"  
/sex="Male"  
/lab\_host="E. Coli strain XL10-Gold, Tn-resistant, F-"  
/clone\_lib="Mouse 10kb plasmid UUGC1M library"  
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 [gil4732114[gb|AF129072.1], a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

**BASE COUNT**  
ORIGIN  
20 a 10 c 8 g 28 t

Query Match 100.0%; Score 10; DB 28; Length 66;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 17 CTTCTCTTTT 26

RESULT 71  
AI442885  
LOCUS  
DEFINITION  
sa28b08.xl1 Gm-cl004 Glycine max cDNA clone GENOME SYSTEMS CLONE ID: Gm-cl004-592 3' similar to TR:024482 024482 SAL13-2.; mRNA sequence.  
AI442885  
VERSION  
AI442885.1 GI:4299305  
KEYWORDS  
EST.

**SOURCE**  
ORGANISM  
Glycine max (soybean)  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae; Glycine. 1 (bases 1 to 67)

**REFERENCE**  
AUTHORS  
Shoemaker R., Keim P., Vodkin L., Erpelding J., Coryell V., Khanna A., Bolla B., Marra M., Hillier L., Kucaba T., Martin J., Beck C., Wylie T., Underwood K., Steptoe M., Theising B., Allen M., Bowers Y., Person B., Swaller T., Gibbons M., Pape D., Harvey N., Schuck R., Ritter E., Kohn S., Shin T., Jackson Y., Cardenas M., McCann R., Waterston R. and Wilson R.

**TITLE**  
Public Soybean EST Project

**JOURNAL**  
Unpublished

**COMMENT**  
Contact: Shoemaker R/Public Soybean EST Project  
Public Soybean EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@watson.wustl.edu  
Trace considered overall poor quality  
Possible reversed clone: similarity on wrong strand This clone is available through: ResGen, Invitrogen Corp. 2130 South Memorial Parkway Huntsville, AL 35801 For further information call: (800)-533-4363 or contact via email: ccu@resgen.com  
Seq primer: -400P from Gibco  
High quality sequence stop: 1  
POLYA=No.

**FEATURES**  
Location/Qualifiers  
1..67  
/organism="Glycine max"  
/mol\_type="mRNA"  
/db\_xref="taxon:3847"  
/clone="GENOME SYSTEMS CLONE ID: Gm-cl004-592"  
/tissue\_type="root"  
/lab\_host="XL10-Gold"  
/clone\_lib="Gm-cl004"  
/note="Vector: pBluescript II XR; Site 1: EcoRI; Site 2: XhoI; Root cDNA. The mRNA was isolated from entire roots of 8 day old 'Williams' seedlings which were propagated on paper towels with distilled water. Stratagene's cDNA Synthesis Kit (catalog #200401) was used to synthesize the cDNA. First-strand synthesis was performed with 5-methyl dCTP, hence the ligated cDNA is hemimethylated. Stratagene's first-strand synthesis primer was used [GAGAGAGAGAGAGAGAGACTAGTCGAG(T)-18]. After second-strand synthesis, the cDNA ends were 'polished' with clone Pfu DNA polymerase, ligated to EcoRI adaptors, and phosphorylated. The XhoI site within the first-strand synthesis primer was restricted by digestion with XhoI; all XhoI sites in the cDNA would be protected by their hemimethylated status. The cDNA constructs were size-fractionated with a 500bp cutoff, using GibcoBRL Life Technologies' cDNA Size Fractionation column. The column eluent was then ligated into Stratagene's pBluescript II XR Predigested vector (pBluescript II SK(+)) that had been digested with EcoRI and XhoI, and phosphorylated). Both the white and blue colonies appear to contain recombinant plasmids with cDNA inserts. Blue colonies 9n=15) have been sequenced, and possess putative cDNA inserts. This library was constructed by Dr. Paul Keim & Virginia H. Coryell, Department of Biology, Box 5640, Northern Arizona University, Flagstaff, AZ 86011, Phone: 520-523-1078 (Dr. Paul Keim), 520-523-1372 (Virginia H. Coryell), Fax: 520-523-7500, email: paul.keim@naui.edu, virginia.coryell@naui.edu"

**BASE COUNT**  
ORIGIN  
18 a 18 c 6 g 24 t 1 others

Query Match 100.0%; Score 10; DB 9; Length 67;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 Db |||||

RESULT 72  
 CD029241  
 LOCUS  
 DEFINITION mgns007xa10f.b Magnaporthe grisea NS Uni-Zap XR Library Magnaporthe grisea cDNA clone mgns007xa10 5', mRNA sequence.  
 ACCESSION CD029241  
 VERSION CD029241  
 KEYWORDS EST.  
 SOURCE CD029241.1 GI:30410697  
 ORGANISM Magnaporthe grisea (anamorph: Pyricularia grisea) Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes; Sordariomycetes incertae sedis; Magnaportheaceae; Magnaporthe. Ebbola, D.J., Yuan, J., Thomas, T.L., Bobrowicz, P., Lu, G., Bhatterai, K. and Dean, R.A.  
 TITLE Expressed sequence tags from the rice blast fungus, Magnaporthe grisea  
 JOURNAL Unpublished  
 COMMENT Contact: Ebbola DJ  
 Department of Plant Pathology & Microbiology  
 Texas A&M University  
 Peterson Bldg, MS2132, College Station, TX 77843-2132, USA  
 Tel: 979 845 4831  
 Fax: 979 845 6483  
 Email: d-ebbola@tamu.edu  
 Chromatogram file of this sequence is available, see contact person

PCR Primers  
 FORWARD: T3 primer  
 BACKWARD: T7 primer  
 Plate: mgns007 row: A column: 10  
 Seq primer: T3.

FEATURES  
 source  
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 Location/Qualifiers  
 /organism="Magnaporthe grisea"  
 /mol\_type="mRNA"  
 /strain="Guy11"  
 /db\_xref="taxon:148305"  
 /clone="mgns007xa10"  
 /sex="Mat1-2 hermaphrodite"  
 /cell\_type="mycelium"  
 /clone\_lib="Magnaporthe grisea NS Uni-Zap XR Library"  
 /note="Vector: pBluescriptSK-; Site 1: EcoRI; Site 2: XhoI ; Unidirectional cloning. EcoRI side has T3 primer and predominantly 5' reads. T7 primer on XhoI side of insert. Nitrogen starvation library. Cells were inoculated into minimal medium and grown for two days with shaking (150 rpm) at room temperature. Culture was harvested, blended, inoculated into minimal medium as above for 24 h. Cells were harvested, washed with water and inoculated into minimal medium base lacking nitrogen source for 6 h. Sequences were processed by one of two methods. Where a full-length alignment to the M. grisea genome sequence was available, the EST sequence was trimmed according to the alignment, otherwise sequence quality was assessed using phredPhrap version 991019 and trimmed according to phd files (0.05) and for vector seqs."

BASE COUNT  
 ORIGIN  
 15 a 18 c 14 g 20 t  
 Query Match 100.0%; Score 10; DB 14; Length 67;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 Db |||||

Db 23 CTTCTCTTTT 32

RESULT 73  
 BM447291/c  
 LOCUS  
 DEFINITION DSA008D10.57203 An expressed sequence tag database for the halotolerant green alga, Dunaliella salina Dunaliella salina cDNA clone DSA008D10 5, mRNA sequence.  
 ACCESSION BM447291  
 VERSION BM447291.1 GI:19852863  
 KEYWORDS EST.  
 SOURCE BM447291.1  
 ORGANISM Dunaliella salina Dunaliella salina Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales; Dunaliellaceae; Dunaliella.  
 REFERENCE 1 (bases 1 to 68)  
 AUTHORS Cushman, J.C.  
 TITLE An expressed sequence tag database for the halotolerant green alga, Dunaliella salina  
 JOURNAL Unpublished  
 COMMENT Contact: Cushman JC  
 Department of Biochemistry  
 University of Nevada  
 MS200, Reno, NV 89557-0014, USA  
 Tel: 775-784-1918  
 Fax: 775-784-1650  
 Email: jcushman@unr.edu  
 PCR Primers  
 FORWARD: T3 20mer  
 BACKWARD: T7 21mer  
 Plate: 008 row: D column: 10  
 Seq primer: T3 20mer  
 High quality sequence stop: 68.

FEATURES  
 source  
 1..68  
 Location/Qualifiers  
 /organism="Dunaliella salina"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:3046"  
 /clone="DSA008D10"  
 /tissue\_type="Cells, which was adapted in 2.5M NaCl via a incremental series from 1.7 to 2.0 to 2.25 to 2.5 M NaCl, were exposed to 3.4 M NaCl for 5 hours"  
 /cell\_type="Green"  
 /clone\_lib="An expressed sequence tag database for the halotolerant green alga, Dunaliella salina"  
 /note="Vector: Lambda Uni-Zap XR, Bluescript SK-; Site 1: EcoRI; Site 2: XhoI; Library construction was performed according to Stratagene's recommended protocol for the Lambda Uni-ZapXR vector and cDNA synthesis kit."

BASE COUNT  
 ORIGIN  
 22 a 13 c 24 g 9 t  
 Query Match 100.0%; Score 10; DB 12; Length 68;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
 Db |||||

RESULT 74  
 AL757142/c  
 LOCUS  
 DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-118D11-012821, genomic survey sequence.  
 ACCESSION AL757142  
 VERSION AL757142.1 GI:21495490  
 KEYWORDS GSS.  
 SOURCE Arabidopsis thaliana (thale cress)  
 ORGANISM Arabidopsis thaliana Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

1 Strizhov, N., Li, Y., Rosso, M., Viehoveer, P., Dekker, K., Saedler, H. and Weisshaar, B.  
A pipeline for automated high-throughput generation of PSTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines  
Unpublished

2 Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weisshaar, B.  
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics  
Unpublished

3 (bases 1 to 68)  
Strizhov, N., Li, Y., Rosso, M. and Weisshaar, B.  
Direct Submission  
Submitted (17-JUN-2002) Weisshaar B., Max-Planck-Institut fuer Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany  
This sequence is recovered from the right border of the T-DNA. It indicates an insertion within the locus defined by clone F27K19. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at:  
<http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES  
source  
1..68  
/organism="Arabidopsis thaliana"  
/mol\_type="genomic DNA"  
/strain="Columbia 0"  
/db\_xref="taxon:3702"  
/clone="GK-118D11-012821"  
/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector PAC161. The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 35 a 8 c 7 g 17 t 1 others  
ORIGIN  
Query Match 100.0%; Score 10; DB 29; Length 68;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 53 CTTCTCTTTT 44

RESULT 75  
AL651716  
LOCUS  
DEFINITION  
AL651716 XGC-gastrula silurana tropicalis cDNA clone TGas036d16 5', mRNA sequence.  
ACCESSION  
AL651716  
VERSION  
AL651716.1 GI:17661928  
KEYWORDS  
EST.  
SOURCE  
Silurana tropicalis (western clawed frog)  
ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Xenopodinae; Silurana.  
1 (bases 1 to 70)  
Huckie, E., Taylor, R., Ashurst, J.L., Zorn, A.M. and Rogers, J.  
Sanger Xenopus tropicalis EST project 2001 (10\_2001)  
Unpublished  
Contact: Huckie E

Sanger Centre  
Hinxton, Cambridgeshire, CB10 1SA, UK  
Email: trop@sanger.ac.uk  
Sanger Xenopus tropicalis EST project 2001  
TROPICALIS\_SEQUENCE\_ID: TGas036d16.sp6  
Sequencing primer: SP6  
This sequence is from a Xenopus Gene Collection (XGC) library constructed by Aaron M. Zorn.

FEATURES  
Location/Qualifiers  
1..70  
/organism="Silurana tropicalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:8364"  
/clone="TGas036d16"  
/dev\_stage="gastrula (stages 10.5-13 mixed)"  
/lab\_host="Escherichia coli XL1-blue"  
/clone\_lib="XGC-gastrula"  
/note="Vector: pCS107; Site 1: EcoRI; Site 2: NotI; cDNA was oligo dt primed from Sug of poly A+ RNA from stages 10-13 gastrulae. EcoRI-NotI cut cDNA was then ligated into pCS107 with EcoRI at the 5' end and NotI at the 3' end."

BASE COUNT 19 a 17 c 15 g 19 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 9; Length 70;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 5 CTTCTCTTTT 14

RESULT 76  
AA255635  
LOCUS  
DEFINITION  
AA255635 70 bp mRNA linear EST 13-AUG-1997  
2931507.r1 NCI CGAP GCBI Homo sapiens cDNA clone IMAGE:668215  
Similar to SW:YKUD\_YEAST P36042 HYPOHETICAL 21.2 KD PROTEIN IN TOR2-PAS1 INTERGENIC REGION.; contains element TARI repetitive element.; mRNA sequence.  
ACCESSION  
AA255635  
VERSION  
AA255635.1 GI:1892570  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens (human)  
ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 70)  
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
Unpublished  
Contact: Robert Strausberg, Ph.D.  
Email: cgaps-remail.nih.gov  
This clone is available royalty-free through LLNL; contact the IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.  
Trace considered overall poor quality  
Possible reversed clone; similarity on wrong strand  
Insert length: 1914 Std Error: 0.00  
Seq primer: -28m13 rev2 ET from Amersham  
High quality sequence stop: 1.  
Location/Qualifiers  
1..70  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:66821"  
/tissue\_type="germinal center B cell"  
/lab\_host="DH10B"  
/clone\_lib="NCI\_CGAP GCBI"  
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA

was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr. David Allman (NCI) and Dr. Gerald Marri (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer (5'-TGTTACCAATCTGAAGTGGAGCGCGCTCAATTTT-3', 15'-TGTTACCAATCTGAAGTGGAGCGCGCTCAATTTT-3'). Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p773 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 0 a 32 c 0 g 38 t  
ORIGIN  
Query Match 100.0%; Score 10; DB 9; Length 70;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 15 CTTCTCTTTT 24

RESULT 77  
H45651/c  
LOCUS  
DEFINITION Yn97d02.s1 Soares adult brain N2B5HB55Y Homo sapiens cDNA clone IMAGE:176355 3' similar to gb|K01562|HUMCRH1 Human Ro RNA (rRNA); ; mRNA sequence.

ACCESSION H45651 70 bp mRNA linear EST 31-JUL-1995  
VERSION H45651.1 GI:921703  
KEYWORDS EST.  
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 70)  
AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lemmon,G., Marra,M., Parsons,J., Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston,R., Williamson,A., Wohlmann,P. and Wilson,R.  
The WashU-Merck EST Project

TITLE Unpublished  
JOURNAL Contact: Wilson RK  
COMMENT Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@watson.wustl.edu  
Insert Size: 979

High quality sequence starts: 1  
High quality sequence stops: 1  
Source: IMAGE Consortium, LLNL  
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.  
Trace considered overall poor quality  
Insert Length: 979 Std Error: 0.00  
Seq primer: SP6  
High quality sequence stop: 1.

FEATURES  
source  
1. .70  
Location/Qualifiers  
/organism="Homo sapiens"  
/mol\_type="rRNA"  
/db\_xref="GDB:3838551"  
/db\_xref="taxon:9606"  
/clone="IMAGE:176355"  
/sex="Male"  
/dev\_stage="55-year old"  
/lab\_host="DH10B (ampicillin resistant)"  
/clone\_lib="Soares adult brain N2B5HB55Y"  
/note="Organ: brain; Vector: p773D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGAGCGCGCTCAATTTT-3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified p773 vector (Pharmacia). Library went through one round of normalization to a Cot = 53. Library constructed by Bento Soares and M. Fatima Bonaldo. The adult brain RNA was provided by Dr. Donald H. Gilden. Tissue was acquired 17-18 hours after death which occurred in consequence of a ruptured aortic aneurysm. RNA was prepared from a pool of tissues representing the following areas of the brain: frontal, parietal, temporal and occipital cortex from the left and right hemispheres, subcortical white matter, basal ganglia, thalamus, cerebellum, midbrain, pons and medulla."

BASE COUNT 27 a 11 c 18 g 14 t  
ORIGIN

Query Match 100.0%; Score 10; DB 14; Length 70;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 13 CTTCTCTTTT 4

RESULT 78

ACCESSION AZ918371/c  
LOCUS AZ918371

DEFINITION 1006004B08.x1 1006 - RescueMu Grid G Zea mays genomic, genomic survey sequence.  
VERSION AZ918371.1 GI:13387655  
KEYWORDS GSS.

SOURCE Zea mays  
ORGANISM Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.  
REFERENCE 1 (bases 1 to 70)  
AUTHORS Walbot V.

TITLE Maize genomic sequences found using engineered RescueMu transposon  
JOURNAL Unpublished  
COMMENT Contact: Walbot V  
Department of Biological Sciences  
Stanford University  
855 California Ave, Palo Alto, CA 94304, USA  
Tel: 650 723 2227  
Fax: 650 725 8221  
Email: walbot@stanford.edu  
Plate: 1006004 row: 36  
Class: transposon-tagged.  
Location/Qualifiers  
1. .70  
/organism="Zea mays"  
/mol\_type="genomic DNA"  
/cultivar="mixed background W23/A188/B73"  
/db\_xref="taxon:4577"  
/tissue\_type="leaf"  
/dev\_stage="adult"  
/lab\_host="DH10B"  
/clone\_lib="1006 - RescueMu Grid G"  
/note="Organ: leaf; Vector: RescueMu (engineered from pBluescript backbone); Site 1: BamHI; Site 2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid G was grown at Stanford in 2000. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B

cells were transformed and then screened on LB plates with ampicillin."

BASE COUNT 23 a 11 c 19 g 17 t

Query Match 100.0%; Score 10; DB 28; Length 70;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 44 CTTCTCTTTT 35

RESULT 79  
D78209/c  
LOCUS D78209 EST from 8p21.3-p22 Homo sapiens cDNA clone B6-1-5, mRNA  
DEFINITION sequence.

ACCESSION D78209.1 GI:2104127

VERSION EST.

KEYWORDS Homo sapiens (human)

SOURCE Homo sapiens

ORGANISM

REFERENCE

AUTHORS

TITLE

Isolation of 45 exon-like fragments from 8p22-->p21.3, a region that is commonly deleted in hepatocellular, colorectal, and non-small cell lung carcinomas

JOURNAL Cytogenet. Cell Genet. 75 (2-3), 190-196 (1996)

MEDLINE 97193198

PUBMED 9040790

COMMENT Contact: Yusuke Nakamura

Institute of Medical Science

University of Tokyo

4-6-1, Shirokanedai, Minato-ku, Tokyo 108, Japan

Tel: 81-3-5449-5372

Fax: 81-3-5449-5433

Email: yusuke@ims.u-tokyo.ac.jp.

Location/Qualifiers

1. .71

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/map="8p21.3-p22"

/clone="B6-1-5"

/clone\_lib="EST from 8p21.3-p22"

29 a 8 c 21 g 13 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 14; Length 71;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 27 CTTCTCTTTT 18

RESULT 80

BZ770025/c

LOCUS SALK\_142959.51.10.x Arabidopsis thaliana cDNA insertion lines

DEFINITION Arabidopsis thaliana genomic clone SALK\_142959.51.10.x, genomic survey sequence.

ACCESSION BZ770025

VERSION BZ770025.1 GI:28943709

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

# REFERENCE AUTHORS

# TITLE JOURNAL COMMENT

# FEATURES source

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 71)

Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab

,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.

,Zimmerman,J. and Ecker,J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@sgalk.edu

This is single pass sequence recovered from the left border of

TDNA. This sequence lies within an annotated intron of At5g01600.

Class: TDNA tagged.

Location/Qualifiers

1. .71

/organism="Arabidopsis thaliana"

/mol\_type="genomic DNA"

/strain="Columbia 0"

/db\_xref="taxon:3702"

/clone="SALK\_142959.51.10.x"

/clone\_lib="Arabidopsis thaliana TDNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines

each of which contains one or more TDNA insertion

elements. The resultant fragment for each line was

directly sequenced to determine the genomic sequence at

the site of insertion. Details of the protocols used can

be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)

28 a 13 c 8 g 22 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 71;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 69 CTTCTCTTTT 60

RESULT 81

AA184862/c

LOCUS AA184862.1 GI:1768508

DEFINITION mu5b11.r1 Soares mouse lymph node NDM1N Mus musculus cDNA clone

IMAGE:642909 5' similar to gb:D13315 LACTOYLGLUTATHIONE LYASE

(HUMAN); mRNA sequence.

72 bp mRNA linear EST 07-JAN-1997

AA184862

VERSION AA184862.1

KEYWORDS Mus musculus (house mouse)

SOURCE Mus musculus

ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

1 (bases 1 to 72)

Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,

Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,

Scheilenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,

Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and

Waterston,R.

The WashU-HMMI Mouse EST Project

Unpublished

Contact: Marra M/Mouse EST Project

WashU-HMMI Mouse EST Project

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810



Email: mouseestowatson.wustl.edu  
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.  
MGI:394901

Trace considered overall poor quality  
Seq primer: -28M13 rev2 from Amersham  
High quality sequence stop: 1.

## FEATURES

source  
Location/Qualifiers  
1..72  
/organism="Mus musculus"  
/mol\_type="mRNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="IMAGE:642909"  
/sex="male"  
/tissue\_type="lymph node"  
/dev\_stages="4 weeks"  
/lab\_host="DH10B"

/note="Organ: lymph node; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not 1; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5', TGTTACCAATCTGAAGTGGAGCGCGCGGATCTTTTTTTTTTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. RNA provided by Dr. Bertrand Jordan. Library constructed and normalized by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 28 a 13 c 17 g 14 t

ORIGIN  
Query Match 100.0%; Score 10; DB 9; Length 72;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
DB 22 CTTCTCTTTT 13

## RESULT 82

LOCUS BX285944 72 bp DNA linear GSS 07-MAR-2003  
DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-386H01-018249, genomic survey sequence.

ACCESSION BX285944  
VERSION BX285944.1 GI:28884940

KEYWORDS GSS  
SOURCE Arabidopsis thaliana (thale cress)

## ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

## REFERENCE

1 Strizhov N., Li, Y., Rosso, M., Viehoever, P., Dekker, K., Saedler, H. and Weisshaar, B.  
A pipeline for automated high-throughput generation of PSTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines  
Unpublished

## JOURNAL

2 Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weisshaar, B.  
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics  
Unpublished

## JOURNAL

3 (bases 1 to 72)

## AUTHORS

Strizhov, N., Rosso, M., Li, Y. and Weisshaar, B.

## TITLE

Submitted (07-MAR-2003) Weisshaar B., Max-Planck-Institut fuer Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany  
This sequence is recovered from the left border of the T-DNA. It indicates an insertion close to or within gene At2g28150. The

sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at:  
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

## FEATURES

source  
Location/Qualifiers  
1..72

/organism="Arabidopsis thaliana"  
/mol\_type="genomic DNA"  
/strain="Columbia 0"  
/db\_xref="taxon:3702"  
/clone="GK-386H01-018249"  
/note="pPCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector pAC161. The lines contain one or more T-DNA from insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 11 a 11 c 16 g 34 t

## ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 72;  
Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
DB 17 CTTCTCTTTT 26

## RESULT 83

LOCUS AA262253/C  
DEFINITION

ACCESSION AA262253

VERSION AA262253.1 GI:1898728

KEYWORDS EST.

SOURCE Homo sapiens (human)

## ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 73)

## REFERENCE

1 Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R.  
WashU-Merck EST Project 1997  
Unpublished

## TITLE

## JOURNAL

## COMMENT

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 1520 Std Error: 0.00

Seq primer: -28M13 rev2 ET from Amersham

High quality sequence stop: 65.

## FEATURES

source  
Location/Qualifiers  
1..73

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:668790"

/tissue\_type="Pooled human melanocyte, fetal heart, and pregnant uterus"



```

/lab host="DH10B"
/clone lib="Soares NHMPU.S1"
/Note="Organ: mixed (see below); Vector: pMT73D-Pac
(Pharmacia) with a modified polylinker; Site_1: Not I;
Site_2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NBHM, pregnant uterus
NbHPU, and fetal heart NbHH19W) were mixed, and ss circles
were made in vitro. Following HAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."
24 a 13 c 15 g 21 t
BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 33 CTTCTCTTTT 24

RESULT 84
AW497651/c
LOCUS
DEFINITION
(SWYD25CAU11A07SK Brugia malayi young adult day 25 cDNA
(SAN99MLW-BmyD25) Brugia malayi cDNA clone SWYD25CAU11A07 5', mRNA
sequence.
ACCESSION
AW497651
VERSION
AW497651.1 GI:7119272
KEYWORDS
EST.
SOURCE
Brugia malayi
ORGANISM
Brugia malayi
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Brugia.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A.
TITLE
Genes expressed in young adult day 25 of Brugia malayi
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
FEATURES
source
1..73
/organism="Brugia malayi"
/mol_type="mRNA"
/db_xref="taxon:6279"
/clone="SWYD25CAU11A07"
/dev_stage="young adult, twenty five days after infection"
/lab_host="XLI-Blue MRF,"
/clone_lib="Brugia malayi young adult day 25 cDNA
(SAN99MLW-BmyD25)"
/notes="Vector: Lambda Uni-Zap XR; Site 1: Eco RI; Site 2:
Xho I; Lymphatic filarial nematode parasite of humans.
mRNA was prepared from young adult worms isolated from
the peritoneal cavity of jirds on day 25 after infection
and converted to double-stranded cDNA using reverse
transcriptase and oligo(dT) followed by RNase H and
pol I. The library has 6.2 x 105 independent recombinants
and the average insert size is approx. 1101bp. The library
was constructed by Michelle Lizotte-Waniewski. The
library is available from Dr. S.A. Williams, email:
genome@neal.smith.edu."
28 a 12 c 21 g 12 t
BASE COUNT

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ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 43 CTTCTCTTTT 34

RESULT 85
AW600116/c
LOCUS
DEFINITION
SWL4CAK10B02SK Brugia malayi L4 cDNA (SAN99MLW-Bml4) Brugia malayi
cDNA clone SWL4CAK10B02 5', mRNA sequence.
ACCESSION
AW600116
VERSION
AW600116.1 GI:7287629
KEYWORDS
EST.
SOURCE
Brugia malayi
ORGANISM
Brugia malayi
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Brugia.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A.
TITLE
Genes expressed in fourth stage larvae of Brugia malayi
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
FEATURES
source
1..73
/organism="Brugia malayi"
/mol_type="mRNA"
/db_xref="taxon:6279"
/clone="SWL4CAK10B02"
/dev_stage="larval stage four"
/lab_host="XLI-Blue MRF,"
/clone_lib="Brugia malayi L4 cDNA (SAN99MLW-Bml4)"
/notes="Vector: Lambda Uni-Zap XR; Site 1: Eco RI; Site 2:
Xho I; Lymphatic filarial nematode parasite of humans.
mRNA was prepared from L4s isolated from the peritoneal
cavity of jirds and converted to double-stranded cDNA
using reverse transcriptase and oligo(dT) followed by
RNase H and DNA pol I. The library has 2.7 x 105
independent recombinants and the average insert size is
approx. 1050bp. The library was constructed by Michelle
Lizotte-Waniewski. The library is available from Dr. S.A.
Williams, email: genome@neal.smith.edu."
30 a 11 c 20 g 12 t
BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 38 CTTCTCTTTT 29

RESULT 86
AW626514/c
LOCUS
DEFINITION
SNOWJ3CAN64C12SK Onchocerca volvulus infective larva cDNA
(SAN94WL-OvJ3) Onchocerca volvulus cDNA clone SNOWJ3CAN64C12 5',
mRNA sequence.

```

ACCESSION AW626514  
 VERSION AW626514.1 GI:7342379  
 KEYWORDS EST.  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 REFERENCE 1 (bases 1 to 73)  
 AUTHORS Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.  
 TITLE Genes expressed in infective third stage larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.  
 FEATURES  
 source  
 1..73  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Sierra Leone"  
 /db\_xref="taxon:6282"  
 /clone="SWOVL3CAN66D01"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Onchocerca volvulus infective larva cDNA (SAM94WL-OvL3)"  
 /note="Vector: lambda UniZap XR; Site\_1: EcoR I; Site\_2: Xho I; Cutaneous filarial nematode parasite of humans. mRNA was prepared from third stage infective larvae of Onchocerca volvulus isolated from mosquitoes 10 days after infection and converted to double stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNase I. The library had 1.8 x 10E5 independent recombinants and average insert size was 900 base pairs. The library was constructed by Wenhong Lu. The library is available from Dr. S.A. Williams, email genome@smith.edu."  
 BASE COUNT 29 a 9 c 22 g 13 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 9; Length 73;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 20 CTTCTCTTTT 11  
 |||||  
 RESULT 87  
 AW626555/c 73 bp mRNA linear EST 30-MAR-2000  
 LOCUS SWOVL3CAN66D01SK Onchocerca volvulus infective larva cDNA  
 DEFINITION (SAM94WL-OvL3) Onchocerca volvulus cDNA clone SWOVL3CAN66D01 5', mRNA sequence.  
 ACCESSION AW626555  
 VERSION AW626555.1 GI:7342420  
 KEYWORDS EST.  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 REFERENCE 1 (bases 1 to 73)  
 AUTHORS Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.  
 TITLE Genes expressed in infective third stage larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology

Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.  
 FEATURES  
 source  
 1..73  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Sierra Leone"  
 /db\_xref="taxon:6282"  
 /clone="SWOVL3CAN66D01"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Onchocerca volvulus infective larva cDNA (SAM94WL-OvL3)"  
 /note="Vector: lambda UniZap XR; Site\_1: EcoR I; Site\_2: Xho I; Cutaneous filarial nematode parasite of humans. mRNA was prepared from third stage infective larvae of Onchocerca volvulus isolated from mosquitoes 10 days after infection and converted to double stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNase I. The library had 1.8 x 10E5 independent recombinants and average insert size was 900 base pairs. The library was constructed by Wenhong Lu. The library is available from Dr. S.A. Williams, email genome@smith.edu."  
 BASE COUNT 26 a 11 c 21 g 15 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 9; Length 73;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 |||||  
 Db 15 CTTCTCTTTT 6  
 |||||  
 RESULT 89  
 AW651817/c 73 bp mRNA linear EST 04-APR-2000  
 LOCUS SWYD25CAUI4C02SK Brugia malayi young adult day 25 cDNA  
 DEFINITION (SAM99MLW-BmyD25) Brugia malayi cDNA clone SWYD25CAUI4C02 5', mRNA sequence.  
 ACCESSION AW651817  
 VERSION AW651817.1 GI:7413075  
 KEYWORDS EST.  
 SOURCE Brugia malayi  
 ORGANISM Brugia malayi  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Brugia.  
 REFERENCE 1 (bases 1 to 73)  
 AUTHORS Genes expressed in young adult day 25 of Brugia malayi  
 TITLE Unpublished  
 JOURNAL Contact: Steven A. Williams  
 COMMENT Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.  
 FEATURES  
 source  
 1..73  
 /organism="Brugia malayi"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:6279"  
 /clone="SWYD25CAUI4C02"  
 /dev\_stage="young adult, twenty five days after infection"  
 /lab\_host="XLI-Blue MRF"

/clone lib="Brugia malayi young adult day 25 cDNA (SAW99MLW-BmYD25)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Lymphatic filarial nematode parasite of humans. mRNA was prepared from young adult worms isolated from the peritoneal cavity of jirds on day 25 after infection and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 6.2 x 10<sup>5</sup> independent recombinants and the average insert size is approx. 1101bp. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr. S.A. Williams, email: genome@neal.smith.edu."

BASE COUNT 29 a 11 c 22 g 11 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 9; Length 73;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 38 CTTCTCTTTT 29  
 |||||

RESULT 89  
 BG310475  
 LOCUS  
 DEFINITION BG310475.1 GI:13112333 73 bp mRNA linear EST 23-FEB-2001  
 (SL96MLW-OvML3) Onchocerca volvulus molting L3 larva cDNA  
 (SL96MLW-OvML3) Onchocerca volvulus cDNA clone SMOV3MCAM55G11 5',  
 mRNA sequence.

ACCESSION BG310475  
 VERSION BG310475.1 GI:13112333  
 KEYWORDS EST.  
 SOURCE Onchocerca volvulus  
 ORGANISM Onchocerca volvulus  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Onchocerca.  
 1 (bases 1 to 73)  
 REFERENCE Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.  
 Genes expressed in molting L3 larvae of Onchocerca volvulus  
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.

FEATURES  
 source  
 1..73  
 /organism="Onchocerca volvulus"  
 /mol\_type="mRNA"  
 /strain="Kumba, Cameroons"  
 /db\_xref="taxon:6282"  
 /clone="SMOV3MCAM55G11"  
 /dev\_stage="molting L3"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Onchocerca volvulus molting L3 larva cDNA (SL96MLW-OvML3)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. Third-stage larvae, L3, were isolated from infected black flies in Cameroon (forest strain). The L3 were cultured in 20% FCS in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in culture. L3 of O. volvulus molt to fourth-stage larvae by day 5 in culture. mRNA was isolated from approximately 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3 in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda

Uni-Zap XR vector and has 1 x 10<sup>5</sup> independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: slustigman@nyc.org)."

BASE COUNT 29 a 10 c 19 g 15 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 10; Length 73;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 35 CTTCTCTTTT 26  
 |||||

RESULT 90  
 AW874933/c  
 LOCUS  
 DEFINITION AW874933 73 bp mRNA linear EST 22-MAY-2000  
 (SWYACAL04D03SK) Brugia malayi young adult cDNA (SAW99MLW-BmYA)  
 Brugia malayi cDNA clone SWYACAL04D03 5', mRNA sequence.

ACCESSION AW874933  
 VERSION AW874933.1 GI:8012644  
 KEYWORDS EST.  
 SOURCE Brugia malayi  
 ORGANISM Brugia malayi  
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;  
 Onchocercidae; Brugia.  
 1 (bases 1 to 73)  
 REFERENCE Williams, S.A.  
 Genes expressed in young adult of Brugia malayi  
 TITLE Genes expressed in young adult of Brugia malayi  
 JOURNAL Unpublished  
 COMMENT Contact: Steven A. Williams  
 Molecular Parasitology  
 Smith College Department of Biological Sciences  
 Department of Biological Sciences, Clark Science Center, Smith  
 College, Northampton, MA, 01063, USA  
 Tel: 4135853826  
 Fax: 4135853786  
 Email: genome@smith.edu  
 Seq primer: pBluescript SK.

FEATURES  
 source  
 1..73  
 /organism="Brugia malayi"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:6279"  
 /clone="SWYACAL04D03"  
 /dev\_stage="young adult"  
 /lab\_host="XLI-Blue MRF"  
 /clone\_lib="Brugia malayi young adult cDNA (SAW99MLW-BmYA)"  
 /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Lymphatic filarial nematode parasite of humans. mRNA was prepared from young adult worms isolated from the peritoneal cavity of jirds and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 6.5 x 10<sup>4</sup> independent recombinants and the average insert size is approx. 800bp. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr. S.A. Williams, email: genome@neal.smith.edu."

BASE COUNT 25 a 10 c 23 g 15 t  
 ORIGIN  
 Query Match 100.0%; Score 10; DB 10; Length 73;  
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTCTCTTTT 10  
 Db 13 CTTCTCTTTT 4  
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```

RESULT 91
BE420470/c
LOCUS      BE420470              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09D05SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09D05 5', mRNA sequence.
ACCESSION  BE420470
VERSION    BE420470.1  GI:9418296
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE  1 (bases 1 to 73)
AUTHORS   Williams, S.A.
TITLE     Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL   Unpublished
COMMENT    Contact: Steven A. Williams
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"
                /clone="SWOVl2CAS09D05"
                /dev_stage="L2"
                /lab_host="XLI-Blue MRF"
                /clone_lib="Onchocerca volvulus L2 larvae cDNA
                (SAW98MLW-OvL2)"
                /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                Xho I; Filarial nematode parasite of humans. mRNA was
                prepared from approximately 9,000 L2s isolated from
                infected mosquitoes from Kumba, Cameroon and converted to
                double-stranded cDNA using reverse transcriptase and
                oligo(dT) followed by RNase H and DNA pol I. The library
                has 7.3 x 10E4 independent recombinants and the average
                insert size is approximately 1kb. The library is
                constructed by Michelle Lizotte-Waniewski. The library is
                available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT 29 a 10 c 19 g 15 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  1  CTTCTCTTTT 10
    |||
Db   19 CTTCTCTTTT 10

RESULT 92
BE420471/c
LOCUS      BE420471              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09D06SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09D06 5', mRNA sequence.
ACCESSION  BE420471
VERSION    BE420471.1  GI:9418297
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE  1 (bases 1 to 73)
AUTHORS   Williams, S.A.

```

```

TITLE     Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL   Unpublished
COMMENT    Contact: Steven A. Williams
            Molecular Parasitology
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"
                /clone="SWOVl2CAS09D06"
                /dev_stage="L2"
                /lab_host="XLI-Blue MRF"
                /clone_lib="Onchocerca volvulus L2 larvae cDNA
                (SAW98MLW-OvL2)"
                /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                Xho I; Filarial nematode parasite of humans. mRNA was
                prepared from approximately 9,000 L2s isolated from
                infected mosquitoes from Kumba, Cameroon and converted to
                double-stranded cDNA using reverse transcriptase and
                oligo(dT) followed by RNase H and DNA pol I. The library
                has 7.3 x 10E4 independent recombinants and the average
                insert size is approximately 1kb. The library is
                constructed by Michelle Lizotte-Waniewski. The library is
                available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT 29 a 10 c 21 g 13 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  1  CTTCTCTTTT 10
    |||
Db   19 CTTCTCTTTT 10

RESULT 93
BE420480/c
LOCUS      BE420480              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09F04SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09F04 5', mRNA sequence.
ACCESSION  BE420480
VERSION    BE420480.1  GI:9418306
KEYWORDS   EST.
SOURCE     Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE  1 (bases 1 to 73)
AUTHORS   Williams, S.A.
TITLE     Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL   Unpublished
COMMENT    Contact: Steven A. Williams
            Molecular Parasitology
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"

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/clone="SWOVL2CAS09F04"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      31 a  12 c  16 g  14 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1  CTTCTCTTTT 10
    |||||
Db  41 CTTCTCTTTT 32

RESULT 94
BE638405/c
LOCUS
DEFINITION
Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ACCESSION
BE638405
VERSION
BE638405.1 GI:9937024
KEYWORDS
EST.
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
TITLE
Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1..73
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS15C01"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      29 a  11 c  19 g  13 t  1 others

```

```

ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1  CTTCTCTTTT 10
    |||||
Db  27 CTTCTCTTTT 18

RESULT 95
BF228818/c
LOCUS
DEFINITION
Onchocerca volvulus L2 larvae cDNA (SAW94WL-OvL3)
ACCESSION
BF228818
VERSION
BF228818.1 GI:11141183
KEYWORDS
EST.
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
TITLE
Genes expressed in infective third stage larvae of Onchocerca
volvulus
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1..73
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Sierra Leone"
/db_xref="taxon:6282"
/clone="SWOVL3CAN76D10"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus infective larva cDNA
(SAW94WL-OvL3)"
/note="Vector: lambda UniZap XR; Site_1: Eco RI; Site_2:
Xho I; Cutaneous filarial nematode parasite of humans.
mRNA was prepared from third stage infective larvae of
Onchocerca volvulus isolated from mosquitoes 10 days after
infection and converted to double stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNAPol I. The library had 1.8 x 10E5 independent
recombinants and average insert size was 900 base pairs.
The library was constructed by Wenhong Lu. The library is
available from Dr. S.A. Williams, email genome@smith.edu."
BASE COUNT      23 a  12 c  22 g  16 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1  CTTCTCTTTT 10
    |||||
Db  11 CTTCTCTTTT 2

RESULT 96
CB886667/c
LOCUS

```

```

DEFINITION      SMOV3MCAM64D11SK Onchocerca volvulus molting L3 larva cDNA
                  (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SMOV3MCAM64D11 5',
                  mRNA sequence.
ACCESSION      CB886667.1 GI:30088462
VERSION        CB886667.1
KEYWORDS       EST.
SOURCE         Onchocerca volvulus
ORGANISM       Onchocerca volvulus
REFERENCE      1 (bases 1 to 73)
AUTHORS        Williams,S.A., Lizotte-Waniewski,M., Laney,S. and Lustigman,S.
TITLE          Genes expressed in molting L3 larvae of Onchocerca volvulus
JOURNAL        Unpublished
COMMENT        Contact: Steven A. Williams
                  Molecular Parasitology
                  Smith College Department of Biological Sciences
                  Department of Biological Sciences, Clark Science Center, Smith
                  College, Northampton, MA, 01063, USA
                  Tel: 4135853826
                  Fax: 4135853786
                  Email: genome@smith.edu
                  Seq primer: pBluescript SK.
FEATURES       Location/Qualifiers
                  1..73
                   /organism="Onchocerca volvulus"
                   /mol_type="mRNA"
                   /strain="Kumba, Cameroons"
                   /db_xref="taxon:6282"
                   /clone="SMOV3MCAM64D11"
                   /dev_stage="molting L3"
                   /lab_host="XLI-Blue MRF"
                   /clone_lib="Onchocerca volvulus molting L3 larva cDNA
                   (SL96MLW-Ovml3)"
                   /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                   Xho I; Filarial nematode parasite of humans. Third-stage
                   larvae, L3, were isolated from infected black flies in
                   Cameroon (forest strain). The L3 were cultured in 20% FCS
                   in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
                   culture. L3 of O. volvulus molt to fourth-stage larvae by
                   day 5 in culture. mRNA was isolated from approximately
                   6000 molting larvae (mL3). 2000 larvae from day 1, 2 or 3
                   in culture, and converted to double-stranded cDNA using
                   reverse transcriptase and oligo(dT) followed by RNase H
                   and DNA pol I. The library was constructed in the lambda
                   Uni-ZAP XR vector and has 1 x 10E6 independent
                   recombinants and the average insert size is ~1200 bp. The
                   library was constructed by Sara Lustigman and Michelle
                   Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.
                   The library is available from Dr. Sara Lustigman (email:
                   slustig@nyc.org)."
BASE COUNT      29 a 10 c 20 g 14 t
ORIGIN
Query Match      100.0%; Score 10; DB 14; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 30 CTTCTCTTTT 21

RESULT 97
BZ660902
LOCUS      BZ660902
DEFINITION      74 bp DNA linear GSS 31-JAN-2003
                  SALK_024363.42.35.x Arabidopsis thaliana TDNA insertion lines
                  Arabidopsis thaliana genomic clone SALK_024363.42.35.x, genomic
                  survey sequence.
ACCESSION      BZ660902
VERSION        BZ660902.1 GI:28174049
KEYWORDS       GSS.
SOURCE         Arabidopsis thaliana (thale cress)

Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Ericaceae; Arabidopsids.
1 (bases 1 to 74)
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: eckersalk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within an annotated exon of At5g56780.
Class: TDNA tagged.
Location/Qualifiers
1..74
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_024363.42.35.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT      15 a 14 c 22 g 22 t
ORIGIN
Query Match      100.0%; Score 10; DB 29; Length 74;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 21 CTTCTCTTTT 30

RESULT 98
CB849383/c
LOCUS      CB849383
DEFINITION      75 bp mRNA linear EST 16-DEC-2002
                  kll1g03.y1 Ascaris suum embryo SL1 TOPO v1 Ascaris suum cDNA 5',
                  mRNA sequence.
ACCESSION      CB849383
VERSION        CB849383.1 GI:27001294
KEYWORDS       EST.
SOURCE         Ascaris suum (pig roundworm)
ORGANISM       Ascaris suum
Eukaryota; Metazoa; Nematoda; Chromadorea; Ascaridida; Ascaridoidea
; Ascarididae; Ascaris.
1 (bases 1 to 75)
McCarter,J., Clifton,S., Chiapelli,B., Pape,D., Martin,J., Wylie,T.,
Dante,M., Marra,M., Hillier,L., Kucaba,T., Theising,B., Bowers,Y.,
Gibbons,M., Ritter,E., Bennett,J., Franklin,C., Tsagarashvili,R.,
Ronko,I., Kennedy,S., Maguire,L., Beck,C., Underwood,K., Steptoe
,M., Allen,M., Person,B., Swaller,T., Harvey,N., Schurk,R., Kohn,S.,
Shin,T., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and
Wilson,R.
The Washington Univ. Nematode EST Project, 1999
Unpublished
Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

```

Tel: 314 286 1800  
Fax: 314 286 1810

Email: estewatson.wustl.edu  
The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Oligo(dT)-SL1 PCR based library. Embryo cDNA PCR products of size >400 nucleotides containing SL1 on the 5' end and oligo(dT) on the 3' end were non-directionally cloned into pCRII-TOPO(Invitrogen) following the Topo TA cloning protocol. 30-60 cell embryo material was provided by Dr. Richard Davis of City University of New York Graduate Center, College of Staten Island, Staten Island, NY (redavis@postbox.csi.cuny.edu).  
Putative full length read  
The vector to vector length is 82  
Seq primer: SL1 primer.

# FEATURES

## Location/Qualifiers

1..75  
/organism="Ascaris suum"  
/mol\_type="mRNA"  
/db\_xref="taxon:6253"  
/dev\_stage="30-60 cell embryo"  
/lab\_host="DH10B"  
/clone\_lib="Ascaris suum embryo SL1 TOPO v1"  
/note="Vector: pCRII-TOPO (Invitrogen); Site 1: EcoRI; Site 2: EcoRI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Oligo(dT)-SL1 PCR based library. Embryo cDNA PCR products of size >400 nucleotides containing SL1 on the 5' end and oligo(dT) on the 3' end were non-directionally cloned into pCRII-TOPO(Invitrogen) following the Topo TA cloning protocol. 30-60 cell embryo material was provided by Dr. Richard Davis of City University of New York Graduate Center, College of Staten Island, Staten Island, NY (redavis@postbox.csi.cuny.edu)."

34 a 10 c 14 g 17 t

## BASE COUNT

34 a 10 c 14 g 17 t

Query Match 100.0%; Score 10; DB 14; Length 75;

Best Local Similarity 100.0%; Pred. No. 1.6e-05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 58 CTCTCTCTTT 49

## RESULT 99

TA77C02P

LOCUS

DEFINITION

TA77C02P 75 bp DNA linear GSS 13-DEC-2000

T. brucei sheared genomic DNA clone 77C02, forward sequence,

genomic survey sequence.

AL460777

AL460777.1 GI:11860526

GSS.

KEYWORDS

SOURCE

ORGANISM

Trypanosoma brucei

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

1 (bases 1 to 75)

Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,

Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,

Melville, S.B., Rajandream, M.A. and Barrell, B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing

project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,

Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and

nh@sanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR),

Rockville, MD. Genomic DNA isolated from a cloned population of

Trypanosoma brucei (FRE927/4 GUTat 10.1) was mechanically sheared

to give a tight size distribution (

4 kb). The v + i method used for the library construction is

described in detail in Smith, H. and Venter, J.C. (Making small

insert libraries for whole genome shotgun sequencing projects. In  
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.  
Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org  
Details of T. brucei sequencing at the Sanger Centre are available  
at [http://www.sanger.ac.uk/Projects/T\\_brucei/](http://www.sanger.ac.uk/Projects/T_brucei/).

## FEATURES

### source

1..75  
/organism="Trypanosoma brucei"  
/mol\_type="genomic DNA"  
/strain="FRE927"  
/db\_xref="taxon:5691"  
/clone="77C02"

13 a 19 c 6 g 37 t

## BASE COUNT

13 a 19 c 6 g 37 t

## ORIGIN

100.0%; Score 10; DB 29; Length 75;

Best Local Similarity 100.0%; Pred. No. 1.6e-05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 19 CTCTCTCTTT 28

## RESULT 100

AA615345/c

LOCUS

DEFINITION

AA615345 76 bp mRNA linear EST 07-OCT-1997

WAGE:1054391 5' similar to SW:TRFE\_RAT P12346 SEROTRANSFERRIN ;

mRNA sequence.

AA615345

AA615345.1 GI:2502573

EST.

KEYWORDS

SOURCE

ORGANISM

Mus musculus

Mus musculus (house mouse)

Eukaryota; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 76)

Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,

Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,

Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,

Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and

Waterston, R.

The WashU-HMMI Mouse EST Project

Unpublished

Contact: Marra M/Mouse EST Project

WashU-HMMI Mouse EST Project

Washington University School

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.

MG1:585967

Trace considered overall poor quality

Possible reversed clone: similarity on wrong strand

Seq primer: -28m13 rev2 ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1..76

/organism="Mus musculus"

/mol\_type="mRNA"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="IMAGE:1054391"

/sex="male"

/tissue\_type="mammary gland"

/dev\_stage="4 weeks"

/lab\_host="DH10B"

/clone\_lib="Soares\_mammary\_gland NbMMG"

/note="Organ: mammary gland; Vector: pT7T3D-Pac (Pharmacia

) with a modified polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5],  
TTCTACCATGTCGAATGGAGCGCCGCAGTCGTATTTTTTTTTTTTTTTTT  
T T 3'; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p7r3 vector.  
The RNA provided by Dr. Minoru Ko, Wayne State Univ. Library constructed and normalized by Bento Soares and M.Fatima Bezade."

22 a	17 c	21 g	16 t
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PAGE CO  
ORIGIN

```
Query Match      100.0%; Score 10; DB 9; Length 76;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

Qy 1 CTTCTCTTTT 10  
Db 34 CTTCTCTTTT 25

Search completed: October 28, 2003, 18:19:28  
Job time : 2140 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:14:39 ; Search time 80 Seconds

(without alignments)

55.173 Million cell updates/sec

Title: US-09-335-032-71

Perfect score: 10

Sequence: 1 cttctctttt 10

Scoring table: OMIGO\_NUC

Gapop 60.0, Gapext 60.0

Searched: 569978 seqs, 220691566 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database : Issued Patents NA.\*  
1: /cgn2\_6/ptodata/2/ina/5A\_COMB.seq.\*  
2: /cgn2\_6/ptodata/2/ina/5B\_COMB.seq.\*  
3: /cgn2\_6/ptodata/2/ina/6A\_COMB.seq.\*  
4: /cgn2\_6/ptodata/2/ina/6B\_COMB.seq.\*  
5: /cgn2\_6/ptodata/2/ina/pctus\_COMB.seq.\*  
6: /cgn2\_6/ptodata/2/ina/backfiles.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	10	100.0	12	2	US-08-173-489C-181
C 2	10	100.0	12	2	US-08-173-489C-246
C 3	10	100.0	20	1	US-08-628-417-3
C 4	10	100.0	20	4	US-09-198-452A-6382
C 5	10	100.0	21	2	US-08-229-528-16
C 6	10	100.0	21	3	US-09-198-484-8
C 7	10	100.0	22	2	US-08-173-489C-65
C 8	10	100.0	24	1	US-08-488-212A-17
C 9	10	100.0	24	2	US-08-320-306-17
C 10	10	100.0	24	2	US-08-488-209B-17
C 11	10	100.0	24	2	US-08-408-011-17
C 12	10	100.0	24	3	US-08-559-205-16
C 13	10	100.0	24	4	US-09-417-722-3
C 14	10	100.0	24	6	5336598-16
C 15	10	100.0	25	1	US-08-628-417-4
C 16	10	100.0	25	6	5217891-9
C 17	10	100.0	27	2	US-08-412-376-23
C 18	10	100.0	27	4	US-08-584-040-7049
C 19	10	100.0	29	4	US-09-270-542-133
C 20	10	100.0	30	4	US-09-270-542-132
C 21	10	100.0	30	4	US-09-270-542-134
C 22	10	100.0	30	4	US-09-270-542-135
C 23	10	100.0	33	3	US-08-961-083-359
C 24	10	100.0	33	4	US-09-536-784-359
C 25	10	100.0	38	4	US-09-371-772B-10906
C 26	10	100.0	39	6	5217891-13
C 27	10	100.0	40	3	US-09-277-016-20
C 1	10	100.0	12	2	Sequence 181, App
C 2	10	100.0	12	2	Sequence 246, App
C 3	10	100.0	20	1	Sequence 3, Appli
C 4	10	100.0	20	4	Sequence 6382, Ap
C 5	10	100.0	21	2	Sequence 16, Appl
C 6	10	100.0	21	3	Sequence 8, Appli
C 7	10	100.0	22	2	Sequence 65, Appl
C 8	10	100.0	24	1	Sequence 17, Appl
C 9	10	100.0	24	2	Sequence 17, Appl
C 10	10	100.0	24	2	Sequence 17, Appl
C 11	10	100.0	24	2	Sequence 17, Appl
C 12	10	100.0	24	3	Sequence 16, Appl
C 13	10	100.0	24	4	Sequence 3, Appli
C 14	10	100.0	24	6	Patent No. 5336598
C 15	10	100.0	25	1	Sequence 4, Appli
C 16	10	100.0	25	6	Patent No. 5217891
C 17	10	100.0	27	2	Sequence 23, Appl
C 18	10	100.0	27	4	Sequence 7049, Ap
C 19	10	100.0	29	4	Sequence 133, App
C 20	10	100.0	30	4	Sequence 132, App
C 21	10	100.0	30	4	Sequence 134, App
C 22	10	100.0	30	4	Sequence 135, App
C 23	10	100.0	33	3	Sequence 359, App
C 24	10	100.0	33	4	Sequence 359, App
C 25	10	100.0	38	4	Sequence 10906, A
C 26	10	100.0	39	6	Patent No. 5217891
C 27	10	100.0	40	3	Sequence 20, Appl

28	10	100.0	42	1	US-07-908-317-33	Sequence 33, Appl
29	10	100.0	42	3	US-09-461-697-227	Sequence 227, App
30	10	100.0	42	5	PCT-US93-06171-33	Sequence 33, Appl
31	10	100.0	43	6	5258302-15	Patent No. 5258302
32	10	100.0	47	4	US-09-671-317-765	Sequence 765, App
33	10	100.0	47	4	US-09-422-978-522	Sequence 522, App
34	10	100.0	47	4	US-09-422-978-1282	Sequence 1282, Ap
35	10	100.0	47	4	US-09-422-978-1546	Sequence 1546, Ap
36	10	100.0	47	4	US-09-422-978-2472	Sequence 2472, Ap
37	10	100.0	47	4	US-09-422-978-3284	Sequence 3284, Ap
38	10	100.0	48	3	US-09-461-697-225	Sequence 225, App
39	10	100.0	50	3	US-08-477-928A-37	Sequence 37, Appl
40	10	100.0	53	3	US-08-910-632-40	Sequence 40, Appl
41	10	100.0	53	3	US-08-910-632-41	Sequence 41, Appl
42	10	100.0	53	3	US-08-805-631A-40	Sequence 40, Appl
43	10	100.0	53	3	US-08-805-631A-41	Sequence 41, Appl
44	10	100.0	53	4	US-09-569-344-40	Sequence 40, Appl
45	10	100.0	53	4	US-09-569-344-41	Sequence 41, Appl
46	10	100.0	54	2	US-08-585-684B-2533	Sequence 2533, Ap
47	10	100.0	54	2	US-08-585-684B-2710	Sequence 2710, Ap
48	10	100.0	54	3	US-09-038-073-2533	Sequence 2533, Ap
49	10	100.0	54	3	US-09-038-073-2710	Sequence 2710, Ap
50	10	100.0	70	4	US-09-446-047A-1	Sequence 1, Appli
51	10	100.0	71	1	US-08-117-374-5	Sequence 5, Appli
52	10	100.0	71	1	US-08-280-263-5	Sequence 5, Appli
53	10	100.0	71	3	US-08-597-325-5	Sequence 5, Appli
54	10	100.0	71	3	US-08-597-325-5	Sequence 5, Appli
55	10	100.0	71	5	PCT-US94-10256-5	Sequence 5, Appli
56	10	100.0	75	1	US-08-117-374-6	Sequence 6, Appli
57	10	100.0	75	1	US-08-280-263-6	Sequence 6, Appli
58	10	100.0	75	3	US-08-597-325-6	Sequence 6, Appli
59	10	100.0	75	3	US-08-597-325-6	Sequence 6, Appli
60	10	100.0	75	5	PCT-US94-10256-6	Sequence 6, Appli
61	10	100.0	78	1	US-08-117-374-19	Sequence 19, Appl
62	10	100.0	78	1	US-08-280-263-19	Sequence 19, Appl
63	10	100.0	78	5	PCT-US94-10256-19	Sequence 19, Appl
64	10	100.0	82	1	US-08-117-374-20	Sequence 20, Appl
65	10	100.0	82	1	US-08-280-263-20	Sequence 20, Appl
66	10	100.0	82	5	PCT-US94-10256-20	Sequence 20, Appl
67	10	100.0	93	3	US-09-174-465D-11	Sequence 11, Appl
68	10	100.0	93	4	US-09-599-564A-11	Sequence 11, Appl
69	10	100.0	96	3	US-09-461-697-223	Sequence 23, App
70	10	100.0	102	4	US-09-144-428-55	Sequence 55, Appl
71	10	100.0	112	3	US-08-932-082-10	Sequence 10, Appl
72	10	100.0	117	3	US-09-461-697-221	Sequence 21, App
73	10	100.0	126	3	US-09-461-697-219	Sequence 219, App
74	10	100.0	132	3	US-09-441-416A-22	Sequence 22, Appl
75	10	100.0	146	3	US-08-477-928A-36	Sequence 36, Appl
76	10	100.0	156	3	US-09-461-697-217	Sequence 217, App
77	10	100.0	174	3	US-09-461-697-215	Sequence 215, App
78	10	100.0	189	3	US-09-461-697-213	Sequence 213, App
79	10	100.0	195	3	US-09-461-697-211	Sequence 211, App
80	10	100.0	198	4	US-09-107-532A-2918	Sequence 2918, Ap
81	10	100.0	201	4	US-09-107-532A-439	Sequence 439, App
82	10	100.0	201	4	US-09-107-532A-3253	Sequence 3253, Ap
83	10	100.0	206	4	US-09-313-294A-7004	Sequence 7004, Ap
84	10	100.0	212	3	US-09-461-697-209	Sequence 209, App
85	10	100.0	222	6	5217891-19	Patent No. 5217891
86	10	100.0	228	4	US-09-016-434-99	Sequence 99, Appl
87	10	100.0	231	3	US-09-461-697-207	Sequence 207, App
88	10	100.0	231	3	US-09-107-532A-2834	Sequence 2834, Ap
89	10	100.0	234	4	US-09-107-532A-724	Sequence 724, App
90	10	100.0	241	4	US-09-397-787-56	Sequence 56, Appl
91	10	100.0	241	4	US-09-389-681-360	Sequence 360, App
92	10	100.0	241	4	US-09-620-405B-360	Sequence 360, App
93	10	100.0	241	4	US-09-433-826B-360	Sequence 360, App
94	10	100.0	241	4	US-09-604-287A-360	Sequence 360, App
95	10	100.0	247	4	US-09-016-434-737	Sequence 737, App
96	10	100.0	251	3	US-09-098-789-9	Sequence 9, Appli
97	10	100.0	256	4	US-09-702-705-1077	Sequence 1077, Ap
98	10	100.0	256	4	US-09-736-457-1077	Sequence 1077, Ap
99	10	100.0	263	4	US-09-313-294A-3143	Sequence 3143, Ap
100	10	100.0	267	4	US-09-134-001C-18	Sequence 18, Appl

C 101	10	100.0	268	4	US-09-313-294A-2020	Sequence 2020, Ap	174	10	100.0	465	6	5496550-9	Patent No. 5496550
C 102	10	100.0	274	4	US-09-313-294A-376	Sequence 376, App	175	10	100.0	466	6	5496550-7	Patent No. 5496550
C 103	10	100.0	276	4	US-09-107-532A-1679	Sequence 1679, Ap	176	10	100.0	475	3	US-08-991-789A-40	Sequence 40, Appl
C 104	10	100.0	278	4	US-09-107-532A-333	Sequence 333, App	177	10	100.0	475	4	US-09-062-451-40	Sequence 5, Appl
C 105	10	100.0	280	2	US-08-731-775-7	Sequence 7, Appl	c 178	10	100.0	475	4	US-09-004-838-5	Sequence 8, Appl
C 106	10	100.0	282	2	US-09-469-697-205	Sequence 205, App	c 179	10	100.0	475	4	US-09-004-838-8	Sequence 40, Appl
C 107	10	100.0	282	4	US-09-016-434-422	Sequence 422, App	c 179	10	100.0	475	4	US-09-598-326-40	Sequence 40, Appl
C 108	10	100.0	285	4	US-09-107-532A-598	Sequence 598, App	181	10	100.0	475	4	US-09-289-198-40	Sequence 27, Appl
C 109	10	100.0	290	3	US-09-203-623-23	Sequence 23, Appl	c 182	10	100.0	477	3	US-08-991-789A-27	Sequence 27, Appl
C 110	10	100.0	290	3	US-09-313-294A-5996	Sequence 5996, Ap	c 183	10	100.0	477	4	US-09-062-451-27	Sequence 27, Appl
C 111	10	100.0	290	4	US-09-523-462-23	Sequence 23, Appl	c 184	10	100.0	477	4	US-08-598-326-27	Sequence 27, Appl
C 112	10	100.0	290	4	US-09-522-980-23	Sequence 23, Appl	c 185	10	100.0	477	4	US-09-328-352-2089	Sequence 2089, Ap
C 113	10	100.0	294	4	US-09-328-352-310	Sequence 310, App	c 186	10	100.0	477	4	US-09-289-198-27	Sequence 27, Appl
C 114	10	100.0	297	4	US-09-313-294A-7153	Sequence 7153, Ap	c 187	10	100.0	480	3	US-08-991-890-3	Sequence 3, Appl
C 115	10	100.0	297	4	US-09-328-352-1780	Sequence 1780, Ap	c 188	10	100.0	484	4	US-09-702-705-645	Sequence 645, App
C 116	10	100.0	300	4	US-09-313-294A-6323	Sequence 6323, Ap	c 189	10	100.0	484	4	US-09-736-457-645	Sequence 645, App
C 117	10	100.0	306	3	US-09-172-108-19	Sequence 19, Appl	c 190	10	100.0	486	1	US-08-828-511-1	Sequence 1, Appl
C 118	10	100.0	306	3	US-09-461-697-203	Sequence 203, App	c 191	10	100.0	486	4	US-09-702-705-877	Sequence 877, App
C 119	10	100.0	309	4	US-09-313-294A-4076	Sequence 4076, Ap	c 192	10	100.0	486	4	US-09-736-457-877	Sequence 877, App
C 120	10	100.0	312	4	US-09-107-532A-2593	Sequence 2593, Ap	c 193	10	100.0	494	3	US-08-477-928A-35	Sequence 35, Appl
C 121	10	100.0	315	3	US-08-617-860B-30	Sequence 30, Appl	c 194	10	100.0	495	4	US-09-328-475C-209	Sequence 209, App
C 122	10	100.0	316	4	US-09-702-705-1553	Sequence 1553, Ap	c 195	10	100.0	498	2	US-08-631-328-54	Sequence 54, Appl
C 123	10	100.0	316	4	US-09-736-457-1553	Sequence 1553, Ap	c 196	10	100.0	498	6	5217891-7	Patent No. 5217891
C 124	10	100.0	318	4	US-09-107-532A-1431	Sequence 1431, Ap	c 197	10	100.0	499	3	US-08-328-111-97	Sequence 97, Appl
C 125	10	100.0	318	4	US-09-107-532A-1846	Sequence 1846, Ap	c 198	10	100.0	499	4	US-09-004-838-9	Sequence 9, Appl
C 126	10	100.0	320	3	US-09-030-607-224	Sequence 224, App	c 199	10	100.0	499	4	US-09-328-475C-211	Sequence 211, App
C 127	10	100.0	320	4	US-09-439-313-224	Sequence 224, App	c 200	10	100.0	501	4	US-09-339-913B-27	Sequence 27, Appl
C 128	10	100.0	320	4	US-09-352-616A-224	Sequence 224, App	c 201	10	100.0	501	4	US-09-339-913B-93	Sequence 93, Appl
C 129	10	100.0	320	4	US-09-232-149A-224	Sequence 224, App	c 202	10	100.0	501	4	US-09-339-913B-98	Sequence 98, Appl
C 130	10	100.0	325	3	US-09-018-584B-26	Sequence 26, Appl	c 203	10	100.0	501	4	US-09-339-904A-87	Sequence 87, Appl
C 131	10	100.0	335	4	US-09-199-542B-110	Sequence 110, App	c 204	10	100.0	501	4	US-09-339-904A-93	Sequence 93, Appl
C 132	10	100.0	337	4	US-09-589-287B-7	Sequence 7, Appl	c 205	10	100.0	501	4	US-09-339-904A-98	Sequence 98, Appl
C 133	10	100.0	337	4	US-09-588-947A-7	Sequence 7, Appl	c 206	10	100.0	501	4	US-08-769-062B-87	Sequence 87, Appl
C 134	10	100.0	339	4	US-09-495-050A-168	Sequence 168, App	c 207	10	100.0	501	4	US-08-769-062B-93	Sequence 93, Appl
C 135	10	100.0	341	1	US-08-322-742-10	Sequence 10, Appl	c 208	10	100.0	501	4	US-08-769-062B-98	Sequence 98, Appl
C 136	10	100.0	342	4	US-09-107-532A-572	Sequence 572, App	c 209	10	100.0	501	4	US-09-344-002B-87	Sequence 87, Appl
C 137	10	100.0	346	4	US-09-171-209-81	Sequence 81, Appl	c 210	10	100.0	501	4	US-09-344-002B-93	Sequence 93, Appl
C 138	10	100.0	354	1	US-07-650-795A-2	Sequence 2, Appl	c 211	10	100.0	501	4	US-09-344-002B-98	Sequence 98, Appl
C 139	10	100.0	357	4	US-09-134-001C-19	Sequence 19, Appl	c 212	10	100.0	501	4	US-09-559-565C-87	Sequence 87, Appl
C 140	10	100.0	360	4	US-09-107-532A-2326	Sequence 2326, Ap	c 213	10	100.0	501	4	US-09-559-565C-93	Sequence 93, Appl
C 141	10	100.0	371	1	US-08-664-596B-25	Sequence 25, Appl	c 214	10	100.0	501	4	US-09-559-565C-98	Sequence 98, Appl
C 142	10	100.0	371	2	US-08-739-775-3	Sequence 3, Appl	c 215	10	100.0	501	4	US-09-693-350-87	Sequence 87, Appl
C 143	10	100.0	372	1	US-08-474-633A-102	Sequence 102, App	c 216	10	100.0	501	4	US-09-693-350-93	Sequence 93, Appl
C 144	10	100.0	372	4	US-08-823-771-102	Sequence 102, App	c 217	10	100.0	501	4	US-09-693-350-98	Sequence 98, Appl
C 145	10	100.0	378	4	US-09-395-996B-3	Sequence 3, Appl	c 218	10	100.0	501	4	US-09-693-389-87	Sequence 87, Appl
C 146	10	100.0	378	4	US-09-395-996B-3	Sequence 3, Appl	c 219	10	100.0	501	4	US-09-693-389-93	Sequence 93, Appl
C 147	10	100.0	378	4	US-09-551-737C-3	Sequence 3, Appl	c 220	10	100.0	501	4	US-09-693-389-98	Sequence 98, Appl
C 148	10	100.0	387	4	US-09-134-001C-253	Sequence 253, App	c 221	10	100.0	510	4	US-09-328-352-2097	Sequence 2097, Ap
C 149	10	100.0	387	4	US-09-134-001C-2156	Sequence 2156, Ap	c 222	10	100.0	515	4	US-09-439-313-472	Sequence 472, App
C 150	10	100.0	393	4	US-09-107-532A-2668	Sequence 2668, Ap	c 223	10	100.0	515	4	US-09-352-616A-472	Sequence 472, App
C 151	10	100.0	394	4	US-09-389-681-448	Sequence 448, App	c 224	10	100.0	517	1	US-08-308-196A-14	Sequence 14, Appl
C 152	10	100.0	394	4	US-09-820-405B-448	Sequence 448, App	c 225	10	100.0	519	4	US-09-205-258-180	Sequence 180, App
C 153	10	100.0	394	4	US-09-433-826B-448	Sequence 448, App	c 226	10	100.0	521	4	US-09-404-879A-128	Sequence 128, App
C 154	10	100.0	394	4	US-09-604-287A-448	Sequence 448, App	c 227	10	100.0	521	4	US-09-338-933-128	Sequence 128, App
C 155	10	100.0	401	4	US-09-643-597-299	Sequence 299, App	c 228	10	100.0	521	4	US-09-215-681-128	Sequence 128, App
C 156	10	100.0	401	4	US-09-408-884A-299	Sequence 299, App	c 229	10	100.0	531	4	US-08-671-548C-41	Sequence 41, Appl
C 157	10	100.0	401	4	US-09-542-615A-299	Sequence 299, App	c 230	10	100.0	531	4	US-08-671-548C-43	Sequence 43, Appl
C 158	10	100.0	401	4	US-09-606-421B-299	Sequence 299, App	c 231	10	100.0	532	4	US-09-679-409-48	Sequence 48, Appl
C 159	10	100.0	407	4	US-09-439-313-402	Sequence 402, App	c 232	10	100.0	533	4	US-09-221-017B-216	Sequence 216, App
C 160	10	100.0	407	4	US-09-352-616A-402	Sequence 402, App	c 233	10	100.0	533	6	5482709-5	Patent No. 5482709
C 161	10	100.0	414	1	US-08-592-126-98	Sequence 98, Appl	c 234	10	100.0	534	4	US-08-671-548C-13	Sequence 13, Appl
C 162	10	100.0	414	4	US-09-168-595-98	Sequence 98, Appl	c 235	10	100.0	537	4	US-08-874-102-42	Sequence 42, Appl
C 163	10	100.0	420	3	US-08-991-890-1	Sequence 1, Appl	c 236	10	100.0	537	4	US-08-874-102-43	Sequence 43, Appl
C 164	10	100.0	421	3	US-09-385-982-49	Sequence 49, Appl	c 237	10	100.0	537	4	US-08-874-102-45	Sequence 45, Appl
C 165	10	100.0	434	4	US-09-280-116-31	Sequence 31, Appl	c 238	10	100.0	537	4	US-08-984-919A-42	Sequence 42, Appl
C 166	10	100.0	438	4	US-09-134-001C-2357	Sequence 2357, Ap	c 239	10	100.0	537	4	US-08-984-919A-43	Sequence 43, Appl
C 167	10	100.0	440	2	US-08-406-057-1	Sequence 1, Appl	c 240	10	100.0	537	4	US-08-984-919A-45	Sequence 45, Appl
C 168	10	100.0	440	3	US-08-958-316-1	Sequence 1, Appl	c 241	10	100.0	542	3	US-08-906-156A-17	Sequence 17, Appl
C 169	10	100.0	451	3	US-09-385-982-55	Sequence 55, Appl	c 242	10	100.0	543	6	5273901-6	Patent No. 5273901
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C 171	10	100.0	456	4	US-09-227-357-110	Sequence 110, App	c 244	10	100.0	544	4	US-09-004-838-137	Sequence 137, App
C 172	10	100.0	456	4	US-09-328-352-2303	Sequence 2303, Ap	c 245	10	100.0	553	4	US-09-171-209-58	Sequence 58, Appl
C 173	10	100.0	460	4	US-09-120-561C-1	Sequence 1, Appl	c 246	10	100.0	553	4		

C 247	10	100.0	555	4	US-09-107-532A-1937	Sequence 1937, Ap	C 320	10	100.0	737	4	US-09-702-705-200	Sequence 200, App
C 248	10	100.0	564	6	5217891-6	Patent No. 5217891	C 321	10	100.0	737	4	US-09-736-457-200	Sequence 5, Appli
C 249	10	100.0	565	3	US-09-328-111-289	Sequence 289, App	C 322	10	100.0	740	2	US-08-739-775-5	Sequence 1, Appli
C 250	10	100.0	573	4	US-08-671-548C-33	Sequence 33, Appl	C 323	10	100.0	757	2	US-08-533-298-1	Sequence 12, Appl
C 251	10	100.0	578	3	US-08-991-789A-222	Sequence 222, App	C 324	10	100.0	762	4	US-09-853-768-12	Sequence 30, Appl
C 252	10	100.0	578	4	US-09-062-451-222	Sequence 222, App	C 325	10	100.0	764	4	US-08-737-109-30	Sequence 47, Appl
C 253	10	100.0	578	4	US-09-598-326-222	Sequence 222, App	C 326	10	100.0	773	4	US-09-227-357-47	Sequence 187, App
C 254	10	100.0	578	4	US-09-289-198-222	Sequence 222, App	C 327	10	100.0	774	3	US-09-461-697-187	Sequence 37, Appl
C 255	10	100.0	582	4	US-08-671-548C-39	Sequence 39, Appl	C 328	10	100.0	776	4	US-09-535-008-37	Sequence 1631, Ap
C 256	10	100.0	582	4	US-08-671-548C-45	Sequence 45, Appl	C 329	10	100.0	780	4	US-09-134-001C-1631	Sequence 5, Appli
C 257	10	100.0	591	4	US-09-702-705-1332	Sequence 1332, Ap	C 330	10	100.0	783	2	US-08-446-922-5	Sequence 1, Appli
C 258	10	100.0	591	4	US-09-702-705-1336	Sequence 1336, Ap	C 331	10	100.0	783	2	US-08-249-189-1	Sequence 1, Appli
C 259	10	100.0	591	4	US-09-736-457-1332	Sequence 1332, Ap	C 332	10	100.0	783	2	US-08-484-624A-1	Sequence 1, Appli
C 260	10	100.0	591	4	US-09-736-457-1336	Sequence 1336, Ap	C 333	10	100.0	783	2	US-08-477-733B-1	Sequence 1, Appli
C 261	10	100.0	594	3	US-09-328-111-155	Sequence 155, App	C 334	10	100.0	783	3	US-09-088-913A-1	Sequence 1, Appli
C 262	10	100.0	599	3	US-09-385-982-446	Sequence 446, App	C 335	10	100.0	783	3	US-08-769-819-1	Sequence 1, Appli
C 263	10	100.0	604	4	US-09-221-017B-1021	Sequence 1021, Ap	C 336	10	100.0	783	3	US-08-770-974-1	Sequence 1, Appli
C 264	10	100.0	607	3	US-09-385-982-239	Sequence 239, App	C 337	10	100.0	783	4	US-08-770-981-1	Sequence 1, Appli
C 265	10	100.0	615	2	US-08-482-842B-1	Sequence 1, Appli	C 338	10	100.0	783	4	US-09-399-106-1	Sequence 1, Appli
C 266	10	100.0	615	2	US-08-482-842B-3	Sequence 3, Appli	C 339	10	100.0	783	5	PCT-US93-10034-5	Sequence 5, Appli
C 267	10	100.0	615	3	US-09-174-465D-1	Sequence 1, Appli	C 340	10	100.0	791	4	US-08-936-165A-132	Sequence 132, App
C 268	10	100.0	615	4	US-09-599-564A-1	Sequence 1, Appli	C 341	10	100.0	791	4	US-09-659-845A-21	Sequence 21, Appl
C 269	10	100.0	621	3	US-09-328-111-204	Sequence 204, App	C 342	10	100.0	795	4	US-09-107-532A-827	Sequence 827, App
C 270	10	100.0	621	3	US-09-328-111-602	Sequence 2838, Ap	C 343	10	100.0	802	3	US-08-441-507-18	Sequence 18, Appl
C 271	10	100.0	621	3	US-09-107-532A-2838	Sequence 93, Appl	C 344	10	100.0	802	4	US-07-969-875A-18	Sequence 121, App
C 272	10	100.0	627	3	US-09-328-111-93	Sequence 104, App	C 345	10	100.0	812	4	US-09-328-475C-121	Sequence 121, App
C 273	10	100.0	628	4	US-09-227-357-104	Sequence 104, App	C 346	10	100.0	813	4	US-09-620-312D-774	Sequence 774, App
C 274	10	100.0	636	4	US-09-107-532A-1567	Sequence 1567, Ap	C 347	10	100.0	816	1	US-09-107-532A-1790	Sequence 1790, Ap
C 275	10	100.0	642	4	US-09-539-333D-114	Sequence 114, App	C 348	10	100.0	818	1	US-08-051-935A-49	Sequence 49, Appl
C 276	10	100.0	642	4	US-09-702-705-77	Sequence 77, Appl	C 349	10	100.0	818	3	US-08-431-055-1	Sequence 1, Appli
C 277	10	100.0	642	4	US-09-736-457-77	Sequence 77, Appl	C 350	10	100.0	818	3	US-08-858-197-1	Sequence 1, Appli
C 278	10	100.0	642	4	US-09-107-532A-1927	Sequence 1927, Ap	C 351	10	100.0	819	3	US-09-461-697-185	Sequence 185, App
C 279	10	100.0	645	4	US-09-328-111-950	Sequence 950, App	C 352	10	100.0	821	3	US-09-361-707-104	Sequence 104, App
C 280	10	100.0	645	4	US-08-961-527-376	Sequence 376, App	C 353	10	100.0	825	4	US-09-280-116-225	Sequence 225, App
C 281	10	100.0	650	4	US-09-404-879A-262	Sequence 262, App	C 354	10	100.0	828	4	US-09-107-532A-1114	Sequence 1114, Ap
C 282	10	100.0	650	4	US-09-338-933-262	Sequence 262, App	C 355	10	100.0	829	3	US-09-961-083-133	Sequence 133, App
C 283	10	100.0	650	4	US-09-215-681-262	Sequence 262, App	C 356	10	100.0	829	4	US-09-556-877-56	Sequence 56, Appl
C 284	10	100.0	660	4	US-09-134-001C-2805	Sequence 2805, Ap	C 357	10	100.0	829	4	US-09-620-412C-56	Sequence 56, Appl
C 285	10	100.0	662	3	US-09-129-030-35	Sequence 35, Appl	C 358	10	100.0	829	4	US-09-410-568-56	Sequence 56, Appl
C 286	10	100.0	669	4	US-09-669-751-79	Sequence 79, Appl	C 359	10	100.0	829	4	US-09-598-419-56	Sequence 56, Appl
C 287	10	100.0	670	3	US-08-791-115B-13	Sequence 13, Appl	C 360	10	100.0	842	4	US-09-536-784-133	Sequence 133, App
C 288	10	100.0	670	4	US-09-140-749-24	Sequence 24, Appl	C 361	10	100.0	842	3	US-08-701-935-2	Sequence 2, Appli
C 289	10	100.0	678	4	US-09-328-352-2378	Sequence 2378, Ap	C 362	10	100.0	846	4	US-09-134-591-2	Sequence 2, Appli
C 290	10	100.0	678	4	US-09-107-532A-3648	Sequence 3648, Ap	C 363	10	100.0	846	4	US-09-328-352-325	Sequence 325, App
C 291	10	100.0	684	4	US-09-328-352-2442	Sequence 2442, Ap	C 364	10	100.0	854	3	US-08-998-416-522	Sequence 522, App
C 292	10	100.0	685	4	US-09-996-243-145	Sequence 145, App	C 365	10	100.0	855	4	US-09-565-423-1	Sequence 1, Appli
C 293	10	100.0	688	1	US-08-090-526-3	Sequence 3, Appli	C 366	10	100.0	865	4	US-09-280-116-59	Sequence 59, Appl
C 294	10	100.0	688	1	US-08-486-715-3	Sequence 3, Appli	C 367	10	100.0	872	3	US-08-943-731-130	Sequence 147, App
C 295	10	100.0	688	1	US-08-486-719-3	Sequence 3, Appli	C 368	10	100.0	872	4	US-09-484-970B-147	Sequence 251, App
C 296	10	100.0	688	1	US-08-271-354-3	Sequence 3, Appli	C 369	10	100.0	876	4	US-09-221-017B-251	Sequence 2, Appli
C 297	10	100.0	688	1	US-08-476-100-3	Sequence 3, Appli	C 370	10	100.0	878	2	US-08-420-786A-2	Sequence 22, Appl
C 298	10	100.0	688	2	US-08-565-861-3	Sequence 3, Appli	C 371	10	100.0	878	2	US-08-484-624A-22	Sequence 22, Appl
C 299	10	100.0	688	3	US-08-475-749-3	Sequence 3, Appli	C 372	10	100.0	878	2	US-08-477-733B-22	Sequence 22, Appl
C 300	10	100.0	688	5	PCT-US94-07658-3	Sequence 3, Appli	C 373	10	100.0	878	3	US-09-088-913A-22	Sequence 22, Appl
C 301	10	100.0	694	2	US-08-537-400-15	Sequence 15, Appl	C 374	10	100.0	878	3	US-09-088-913A-22	Sequence 22, Appl
C 302	10	100.0	694	2	US-08-706-702-17	Sequence 17, Appl	C 375	10	100.0	878	3	US-08-769-819-22	Sequence 22, Appl
C 303	10	100.0	694	3	US-08-706-706-17	Sequence 17, Appl	C 376	10	100.0	878	4	US-08-770-974-22	Sequence 22, Appl
C 304	10	100.0	694	4	US-09-238-471-17	Sequence 17, Appl	C 377	10	100.0	878	4	US-08-770-981-22	Sequence 22, Appl
C 305	10	100.0	696	3	US-09-461-697-193	Sequence 193, App	C 378	10	100.0	882	4	US-09-399-106-22	Sequence 22, Appl
C 306	10	100.0	699	3	US-09-461-697-193	Sequence 193, App	C 379	10	100.0	882	3	US-09-068-140A-1	Sequence 1, Appli
C 307	10	100.0	705	4	US-09-107-532A-2425	Sequence 2425, Ap	C 380	10	100.0	903	4	US-09-134-001C-1260	Sequence 1260, Ap
C 308	10	100.0	711	4	US-09-662-250A-13	Sequence 13, Appl	C 381	10	100.0	903	4	US-09-589-287B-18	Sequence 18, Appl
C 309	10	100.0	714	4	US-09-107-532A-3372	Sequence 3372, Ap	C 382	10	100.0	903	4	US-09-588-947A-18	Sequence 18, Appl
C 310	10	100.0	717	3	US-09-461-697-189	Sequence 189, App	C 383	10	100.0	906	4	US-09-134-001C-2072	Sequence 2072, Ap
C 311	10	100.0	718	3	US-08-998-416-961	Sequence 961, App	C 384	10	100.0	907	1	US-08-664-596B-23	Sequence 23, App
C 312	10	100.0	720	1	US-08-153-848-35	Sequence 35, Appl	C 385	10	100.0	907	2	US-09-035-648-19	Sequence 19, Appl
C 313	10	100.0	720	2	US-08-722-349-4	Sequence 4, Appli	C 386	10	100.0	909	3	US-08-739-775-1	Sequence 1, Appli
C 314	10	100.0	720	2	US-09-204-328-4	Sequence 4, Appli	C 387	10	100.0	909	3	US-09-001-951-19	Sequence 19, Appl
C 315	10	100.0	720	3	US-09-299-843A-35	Sequence 35, Appl	C 388	10	100.0	909	4	US-08-818-829-19	Sequence 19, Appl
C 316	10	100.0	720	4	US-09-088-337B-35	Sequence 35, Appl	C 389	10	100.0	921	4	US-09-107-532A-896	Sequence 896, App
C 317	10	100.0	720	4	US-09-107-532A-3104	Sequence 3104, Ap	C 390	10	100.0	927	4	US-09-328-352-1986	Sequence 1986, Ap
C 318	10	100.0	720	5	PCT-US93-11153-35	Sequence 35, Appl	C 391	10	100.0	929	2	US-08-586-676B-9	Sequence 9, Appli
C 319	10	100.0	723	2	US-08-680-395-2	Sequence 2, Appli	C 392	10	100.0	930	4	US-09-107-532A-2737	Sequence 2737, Ap

C 393	10	100.0	934	3	US-09-174-465D-4	Sequence 4, Appli	Sequence 4, Appli
C 394	10	100.0	934	4	US-09-599-564A-4	Sequence 4, Appli	Sequence 4, Appli
C 395	10	100.0	945	4	US-09-328-352-2968	Sequence 2968, Ap	Sequence 2968, Ap
C 396	10	100.0	951	4	US-09-556-877-118	Sequence 118, App	Sequence 118, App
C 397	10	100.0	951	4	US-09-620-412C-118	Sequence 118, App	Sequence 118, App
C 398	10	100.0	951	4	US-09-598-419-118	Sequence 118, App	Sequence 118, App
C 399	10	100.0	957	1	US-08-381-280-2	Sequence 2, Appli	Sequence 2, Appli
C 400	10	100.0	957	1	US-08-381-280-7	Sequence 7, Appli	Sequence 7, Appli
C 401	10	100.0	957	1	US-08-381-280-16	Sequence 16, Appl	Sequence 16, Appl
C 402	10	100.0	957	2	US-08-445-533-2	Sequence 2, Appli	Sequence 2, Appli
C 403	10	100.0	957	2	US-08-445-533-7	Sequence 7, Appli	Sequence 7, Appli
C 404	10	100.0	957	2	US-08-445-533-16	Sequence 16, Appl	Sequence 16, Appl
C 405	10	100.0	957	3	US-09-052-085-2	Sequence 2, Appli	Sequence 2, Appli
C 406	10	100.0	957	3	US-09-052-085-7	Sequence 7, Appli	Sequence 7, Appli
C 407	10	100.0	957	3	US-09-052-085-16	Sequence 16, Appl	Sequence 16, Appl
C 408	10	100.0	957	4	US-08-858-207A-104	Sequence 104, App	Sequence 104, App
C 409	10	100.0	960	4	US-09-134-001C-2618	Sequence 2618, Ap	Sequence 2618, Ap
C 410	10	100.0	975	6	5340934-10	Patent No. 5340934	Patent No. 5340934
C 411	10	100.0	977	6	5340934-9	Sequence 676, App	Sequence 676, App
C 412	10	100.0	978	4	US-09-134-001C-676	Sequence 9, Appli	Sequence 9, Appli
C 413	10	100.0	984	3	US-08-748-506-9	Sequence 15, Appl	Sequence 15, Appl
C 414	10	100.0	985	4	US-07-145-002B-15	Patent No. 5217891	Patent No. 5217891
C 415	10	100.0	987	6	5217891-3	Sequence 283, App	Sequence 283, App
C 416	10	100.0	989	4	US-09-671-317-283	Sequence 636, App	Sequence 636, App
C 417	10	100.0	1000	4	US-09-641-638-636	Sequence 637, App	Sequence 637, App
C 418	10	100.0	1000	4	US-09-641-638-637	Sequence 168, App	Sequence 168, App
C 419	10	100.0	1001	4	US-09-641-638-168	Sequence 169, App	Sequence 169, App
C 420	10	100.0	1001	4	US-09-641-638-169	Sequence 555, App	Sequence 555, App
C 421	10	100.0	1001	4	US-09-641-638-555	Sequence 174, App	Sequence 174, App
C 422	10	100.0	1001	4	US-09-671-317-174	Sequence 272, App	Sequence 272, App
C 423	10	100.0	1001	4	US-09-671-317-272	Sequence 273, App	Sequence 273, App
C 424	10	100.0	1001	4	US-09-671-317-273	Sequence 309, App	Sequence 309, App
C 425	10	100.0	1001	4	US-09-671-317-309	Sequence 333, App	Sequence 333, App
C 426	10	100.0	1001	4	US-09-671-317-333	Sequence 334, App	Sequence 334, App
C 427	10	100.0	1001	4	US-09-671-317-334	Sequence 352, App	Sequence 352, App
C 428	10	100.0	1001	4	US-09-671-317-352	Sequence 353, App	Sequence 353, App
C 429	10	100.0	1001	4	US-09-671-317-353	Sequence 401, App	Sequence 401, App
C 430	10	100.0	1001	4	US-09-671-317-401	Sequence 402, App	Sequence 402, App
C 431	10	100.0	1001	4	US-09-671-317-402	Sequence 453, App	Sequence 453, App
C 432	10	100.0	1001	4	US-09-671-317-453	Sequence 69, Appl	Sequence 69, Appl
C 433	10	100.0	1004	3	US-08-714-918-69	Sequence 69, Appl	Sequence 69, Appl
C 434	10	100.0	1004	3	US-09-265-315-69	Sequence 69, Appl	Sequence 69, Appl
C 435	10	100.0	1004	3	US-09-265-315-69	Sequence 69, Appl	Sequence 69, Appl
C 436	10	100.0	1004	3	US-09-266-417-69	Sequence 69, Appl	Sequence 69, Appl
C 437	10	100.0	1011	4	US-09-684-938-178	Sequence 178, App	Sequence 178, App
C 438	10	100.0	1011	4	US-09-308-825A-178	Sequence 61, Appl	Sequence 61, Appl
C 439	10	100.0	1021	4	US-09-465-558-61	Sequence 90, Appl	Sequence 90, Appl
C 440	10	100.0	1024	4	US-09-328-475C-90	Sequence 2, Appli	Sequence 2, Appli
C 441	10	100.0	1026	1	US-08-423-568-2	Sequence 1353, Ap	Sequence 1353, Ap
C 442	10	100.0	1029	4	US-09-134-001C-1353	Sequence 16, Appl	Sequence 16, Appl
C 443	10	100.0	1040	4	US-09-288-143-16	Sequence 31, Appl	Sequence 31, Appl
C 444	10	100.0	1046	1	US-08-362-670B-31	Sequence 31, Appl	Sequence 31, Appl
C 445	10	100.0	1046	1	US-08-333-576C-31	Sequence 31, Appl	Sequence 31, Appl
C 446	10	100.0	1046	1	US-08-808-324-31	Sequence 31, Appl	Sequence 31, Appl
C 447	10	100.0	1046	5	PCT-US94-14030A-31	Sequence 31, Appl	Sequence 31, Appl
C 448	10	100.0	1051	2	US-08-865-273-1	Sequence 1, Appli	Sequence 1, Appli
C 449	10	100.0	1051	4	US-09-385-174-1	Sequence 1, Appli	Sequence 1, Appli
C 450	10	100.0	1051	4	US-08-715-252-1	Sequence 1, Appli	Sequence 1, Appli
C 451	10	100.0	1082	2	US-08-453-051-3	Sequence 3, Appli	Sequence 3, Appli
C 452	10	100.0	1100	4	US-09-589-287B-1	Sequence 1, Appli	Sequence 1, Appli
C 453	10	100.0	1100	4	US-09-588-947A-1	Sequence 13, Appl	Sequence 13, Appl
C 454	10	100.0	1101	4	US-08-984-919A-13	Sequence 14, Appl	Sequence 14, Appl
C 455	10	100.0	1101	4	US-08-984-919A-14	Sequence 1, Appli	Sequence 1, Appli
C 456	10	100.0	1105	3	US-08-837-317-1	Sequence 1, Appli	Sequence 1, Appli
C 457	10	100.0	1105	4	US-09-573-885A-1	Sequence 13, Appl	Sequence 13, Appl
C 458	10	100.0	1107	3	US-08-781-420-13	Sequence 14, Appl	Sequence 14, Appl
C 459	10	100.0	1107	3	US-08-781-420-14	Sequence 13, Appl	Sequence 13, Appl
C 460	10	100.0	1107	4	US-08-874-102-13	Sequence 14, Appl	Sequence 14, Appl
C 461	10	100.0	1107	4	US-08-874-102-14	Sequence 13, Appl	Sequence 13, Appl
C 462	10	100.0	1107	4	US-09-006-595A-13	Sequence 14, Appl	Sequence 14, Appl
C 463	10	100.0	1107	4	US-09-006-595A-14	Sequence 31, Appl	Sequence 31, Appl
C 464	10	100.0	1109	4	US-07-145-002B-31	Sequence 1614, Ap	Sequence 1614, Ap
C 465	10	100.0	1113	4	US-09-107-532A-1614		

Sequence 1192, Ap

Sequence 11, Appl

Sequence 102, App

Sequence 2, Appli

Sequence 6, Appli

Sequence 19, Appl

Sequence 23, Appl

Sequence 26, Appl

Sequence 29, Appl

Sequence 610, App

Sequence 3, Appli

Sequence 1, Appli

Sequence 15, Appl

Sequence 1000, Ap

Sequence 705, App

Sequence 8, Appli

Sequence 3, Appli

Sequence 2399, Ap

Sequence 35, Appl

Sequence 36, Appl

Sequence 36, Appl

Sequence 3, Appli

Sequence 22, Appl

Sequence 22, Appl

Sequence 32, Appl

Sequence 4, Appli

Sequence 169, App

Sequence 169, App

Sequence 169, App

Sequence 1, Appli

Sequence 1039, Ap

Sequence 2488, Ap

Sequence 569, App

US-08-173-489C-181/c

Sequence 181, Application US/08173489C

Patent No. 5861344

GENERAL INFORMATION:

APPLICANT: WANG, C.-G.

APPLICANT: HEPBURN, A. G.

TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA

TITLE OF INVENTION: TRIPLE-STRAND FORMATION.

NUMBER OF SEQUENCES: 365

CORRESPONDENCE ADDRESS:

ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,

STREET: 510 EAST 73RD STREET,

CITY: NEW YORK

STATE: NEW YORK

COUNTRY: USA

ZIP: 10023.

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44mb storage

COMPUTER: IBM PC/XT/AT

OPERATING SYSTEM: MS-DOS version 6.2

SOFTWARE: Wordperfect Version 5.1

CURRENT APPLICATION DATA: US/08/173,489C

APPLICATION NUMBER: 22 DEC 1993

FILING DATE: 22 DEC 1993

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/968,436

FILING DATE: 29 OCT 1992

ATTORNEY/AGENT INFORMATION:

NAME: Handelman, Joseph H.

REGISTRATION NUMBER: 26,179

REFERENCE/DOCKET NUMBER: U9518-6

RESULT 1

US-08-173-489C-181/c

Sequence 181, Application US/08173489C

Patent No. 5861344

GENERAL INFORMATION:

APPLICANT: WANG, C.-G.

APPLICANT: HEPBURN, A. G.

TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA

TITLE OF INVENTION: TRIPLE-STRAND FORMATION.

NUMBER OF SEQUENCES: 365

CORRESPONDENCE ADDRESS:

ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,

STREET: 510 EAST 73RD STREET,

CITY: NEW YORK

STATE: NEW YORK

COUNTRY: USA

ZIP: 10023.

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44mb storage

COMPUTER: IBM PC/XT/AT

OPERATING SYSTEM: MS-DOS version 6.2

SOFTWARE: Wordperfect Version 5.1

CURRENT APPLICATION DATA: US/08/173,489C

APPLICATION NUMBER: 22 DEC 1993

FILING DATE: 22 DEC 1993

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/968,436

FILING DATE: 29 OCT 1992

ATTORNEY/AGENT INFORMATION:

NAME: Handelman, Joseph H.

REGISTRATION NUMBER: 26,179

REFERENCE/DOCKET NUMBER: U9518-6

```

;
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (attorney) (212) 708-1880
; TELEFAX: (attorney) (212) 246-8959
; INFORMATION FOR SEQ ID NO: 181:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 12 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double stranded
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; DESCRIPTION: hepatitis B virus adw2 isolate,
; DESCRIPTION: nucleotides 807 to 818
; HYPOTHETICAL: no
; ANTI-SENSE: no
; ORGANISM: Hepatitis B virus
; INDIVIDUAL ISOLATE: adw2
; PUBLICATION INFORMATION:
; AUTHORS: Valenzuela, P, Quiroga, M, Zaldivar, J,
; AUTHORS: Gray, P, Ruter, W J.
; TITLE: The nucleotide sequence of
; TITLE: the Hepatitis B viral genome and the
; TITLE: identification of the major viral genes
; JOURNAL: In "Animal Virus Genetics", Fields, B N,
; JOURNAL: Jaenisch, R, Fox C F eds
; VOLUME:
; PAGES: 57-70
; DATE: 1980
; RELEVANT RESIDUES IN SEQ ID NO: 181 :FROM 1 TO 12
;
US-08-173-489C-181

Query Match 100.0%; Score 10; DB 2; Length 12;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 2
US-08-173-489C-246
; Sequence 246, Application US/08173489C
; Patent No. 5861244
; GENERAL INFORMATION:
; APPLICANT: WANG, C. -G.
; APPLICANT: HEPBURN, A. G.
; TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA
; TITLE OF INVENTION: TRIPLE-STRAND FORMATION.
; NUMBER OF SEQUENCES: 365
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,
; STREET: 510 EAST 73RD STREET,
; CITY: NEW YORK
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 10021.
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch, 1.44Mb storage
; COMPUTER: IBM PC/XT/AT
; OPERATING SYSTEM: MS-DOS version 6.2
; SOFTWARE: Wordperfect Version 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/173,489C
; FILING DATE: 22 DEC 1993
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/969,436
; FILING DATE: 29 OCT 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Handelman, Joseph H.
; REGISTRATION NUMBER: 26,179
; REFERENCE/DOCKET NUMBER: U9518-6

;
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (attorney) (212) 708-1880
; TELEFAX: (attorney) (212) 246-8959
; INFORMATION FOR SEQ ID NO: 246:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 12 bases
; TYPE: nucleic acid
; STRANDEDNESS: single stranded
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: third strand derived from M. luteus
; DESCRIPTION: 23s region in Seq ID No. 5861244245
; HYPOTHETICAL: yes
; ANTI-SENSE: no
; PUBLICATION INFORMATION:
; RELEVANT RESIDUES IN SEQ ID NO: 246 :FROM 1 TO 12
;
US-08-173-489C-246

Query Match 100.0%; Score 10; DB 2; Length 12;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 3 CTTCTCTTTT 12

RESULT 3
US-08-628-417-3
; Sequence 3, Application US/08628417
; Patent No. 5627054
; GENERAL INFORMATION:
; APPLICANT: GILLESPIE, DAVID
; TITLE OF INVENTION: COMPETITOR PRIMER ASYMMETRIC
; TITLE OF INVENTION: POLYMERASE CHAIN REACTION
; NUMBER OF SEQUENCES: 7
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: U.S. ARMY CHEMICAL AND BIOLOGICAL
; ADDRESSEE: DEFENSE COMMAND
; STREET: OFFICE OF THE CHIEF COUNSEL (AMSCB-GC)
; CITY: ABERDEEN PROVING GROUND
; STATE: MARYLAND
; COUNTRY: USA
; ZIP: 21010-5423
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/628,417
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: BIFFONI, ULYSSES J
; REGISTRATION NUMBER: 39,908
; REFERENCE/DOCKET NUMBER: DAM 398-94
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 410-671-1158
; TELEFAX: 410-671-2534
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 bases
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: oligodeoxynucleotide
; HYPOTHETICAL: NO
; ANTI-SENSE: YES
; ORIGINAL SOURCE:
; ORGANISM: Staphylococcus aureus
;
US-08-628-417-3
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Query Match      100.0%; Score 10; DB 1; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      8 CTTCTCTTTT 17

RESULT 4
US-09-198-452A-6382/c
; Sequence 6382, Application US/09198452A
; Patent No. 6559294
; GENERAL INFORMATION:
; APPLICANT: Griffais, R.
; TITLE OF INVENTION: Chlamydia pneumoniae genomic sequence and polypeptides, fragments
; TITLE OF INVENTION: thereof and uses thereof, in particular for the diagnosis, prevention
; TITLE OF INVENTION: and treatment of infection
; FILE REFERENCE: 9710-003-999
; CURRENT APPLICATION NUMBER: US/09/198,452A
; CURRENT FILING DATE: 1998-11-24
; NUMBER OF SEQ ID NOS: 6949
; SEQ ID NO 6382
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia pneumoniae
US-09-198-452A-6382

Query Match      100.0%; Score 10; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      17 CTTCTCTTTT 8

RESULT 5
US-08-229-528-16/c
; Sequence 16, Application US/08229528
; Patent No. 5837447
; GENERAL INFORMATION:
; APPLICANT: GORSKI, Jack
; TITLE OF INVENTION: MONITORING AN IMMUNE RESPONSE BY ANALYSIS OF AMPLIFIED IMMUNO
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Iardner
; STREET: P. O. Box 1497
; CITY: Madison
; STATE: Wisconsin
; COUNTRY: USA
; ZIP: 53701-1497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.44 Mb storage
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: MS-DOS 3.3
; SOFTWARE: Wordperfect, Version 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/229,528
; FILING DATE: 18-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/868,569
; FILING DATE: 15-APR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Scanlon, William J.
; REGISTRATION NUMBER: 30,136
; REFERENCE/DOCKET NUMBER: 30383/133
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (608) 258-4284
; TELEFAX: (608) 258-4258
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
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; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: Other nucleic acid;
; DESCRIPTION: Synthetic DNA oligonucleotide
US-08-229-528-16

Query Match      100.0%; Score 10; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      18 CTTCTCTTTT 9

RESULT 6
US-09-198-484-8/c
; Sequence 8, Application US/09198484
; Patent No. 6162435
; GENERAL INFORMATION:
; APPLICANT: Minion, F. Chris
; APPLICANT: Hsu, Tsungda
; TITLE OF INVENTION: RECOMBINANT MYCOPLASMA HYOPNEUMONIAE VACCINE
; FILE REFERENCE: I9000.028/P028
; CURRENT APPLICATION NUMBER: US/09/198,484
; CURRENT FILING DATE: 1998-11-24
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 21
; TYPE: DNA
; ORGANISM: primer
US-09-198-484-8

Query Match      100.0%; Score 10; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      15 CTTCTCTTTT 6

RESULT 7
US-08-173-489C-65
; Sequence 65, Application US/08173489C
; Patent No. 5861244
; GENERAL INFORMATION:
; APPLICANT: WANG, C. -G.
; APPLICANT: HEPBURN, A. G.
; TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA
; TITLE OF INVENTION: TRIPLE-STRAND FORMATION.
; NUMBER OF SEQUENCES: 365
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,
; STREET: 510 EAST 73RD STREET,
; CITY: NEW YORK
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 10021
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch, 1.44Mb storage
; COMPUTER: IBM PC/XT/AT
; OPERATING SYSTEM: MS-DOS version 6.2
; SOFTWARE: Wordperfect Version 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/173,489C
; FILING DATE: 22 DEC 1993
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/968,436
; FILING DATE: 29 OCT 1992
```



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; FILING DATE: No. 5891623ember 9, 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Thomas E. Popovich
; REGISTRATION NUMBER: 30099
; REFERENCE/DOCKET NUMBER: 3678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (612) 334-8991
; TELEFAX: (612) 334-8994
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 bases
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; MOLECULE TYPE: Other nucleic acid
; MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
; MOLECULE TYPE: Vb region)
; HYPOTHETICAL: No
; ORIGINAL SOURCE: Synthesized using
; PUBLICATION INFORMATION:
; AUTHORS: Imberti, Luisa; Sottini,
; AUTHORS: Alessandra; Bettinardi, Alessandria; Puoti, Massimo; Primi,
; AUTHORS: Daniele
; TITLE: Selective Depletion in HIV Infection
; TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
; JOURNAL: Science
; VOLUME: 254
; ISSUE: 5033
; PAGES: 860-862
; PUBLICATION DATE: No. 5891623ember 8, 1991
; US-08-320-306-17

```

```

Query Match 100.0%; Score 10; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

```

```

RESULT 10
US-08-488-209B-17/c
; Sequence 17, Application US/08488209B
; Patent No. 5925513
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft Word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488.209B
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485
; FILING DATE: No. 5925513ember 9, 1992

```

```

; ATTORNEY/AGENT INFORMATION:
; NAME: Thomas E. Popovich
; REGISTRATION NUMBER: 30099
; REFERENCE/DOCKET NUMBER: 3678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (612) 334-8991
; TELEFAX: (612) 334-8994
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 bases
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; MOLECULE TYPE: Other nucleic acid
; MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
; MOLECULE TYPE: Vb region)
; HYPOTHETICAL: No
; ORIGINAL SOURCE: Synthesized using
; PUBLICATION INFORMATION:
; AUTHORS: Imberti, Luisa; Sottini,
; AUTHORS: Alessandra; Bettinardi, Alessandria; Puoti, Massimo; Primi,
; AUTHORS: Daniele
; TITLE: Selective Depletion in HIV Infection
; TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
; JOURNAL: Science
; VOLUME: 254
; ISSUE: 5033
; PAGES: 860-862
; PUBLICATION DATE: No. 5925513ember 8, 1991
; US-08-488-209B-17

Query Match 100.0%; Score 10; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

```

```

RESULT 11
US-08-408-011-17/c
; Sequence 17, Application US/08408011
; Patent No. 5928642
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft Word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/408.011
; FILING DATE: 18-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485
; FILING DATE: No. 5928642ember 9, 1992
; ATTORNEY/AGENT INFORMATION:

```



NAME: Thomas E. Popovich  
REGISTRATION NUMBER: 30099  
REFERENCE/DOCKET NUMBER: 3678  
TELEPHONE: (612) 334-8991  
TELEFAX: (612) 334-8994  
INFORMATION FOR SEQ ID NO: 17:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 24 bases  
TYPE: Nucleic Acid  
STRANDEDNESS: Single  
TOPOLOGY: Linear  
MOLECULE TYPE: Other nucleic acid  
MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor  
MOLECULE TYPE: Vb region)  
HYPOTHETICAL: No  
ORIGINAL SOURCE: Synthesized using  
PUBLICATION INFORMATION:  
AUTHORS: Imberti, Luisa; Sottini,  
AUTHORS: Alessandra; Bettinardi, Alessandra; Puoti, Massimo; Primi,  
AUTHORS: Daniele  
TITLE: Selective Depletion in HIV Infection  
TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences  
JOURNAL: Science  
VOLUME: 254  
ISSUE: 5033  
PAGES: 860-862  
PUBLICATION DATE: No. 5928642ember 8, 1991  
US-08-408-011-17

Query Match 100.0%; Score 10; DB 2; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 18 CTTCTCTTTT 9

RESULT 12  
US-08-559-205-16/c  
Sequence 16, Application US/08559205  
Patent No. 6087096  
GENERAL INFORMATION:  
APPLICANT: Dau, Peter C.  
APPLICANT: Liu, Debang  
TITLE OF INVENTION: Method of Intrafamily Fragment Analysis of the T  
TITLE OF INVENTION: Cell Receptor ' and Chain CDR3 Regions  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun  
STREET: 6300 Sears Tower, 233 South Wacker Drive  
CITY: Chicago  
STATE: Illinois  
COUNTRY: United States of America  
ZIP: 60606-6402  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/559,205  
FILING DATE:  
CLASSIFICATION: 436  
ATTORNEY/AGENT INFORMATION:  
NAME: Gass, David A.  
REGISTRATION NUMBER: 38,153  
REFERENCE/DOCKET NUMBER: 28721/32972  
TELEPHONE: 312/474-6300  
TELEFAX: 312/474-0448

TELEX: 25-3856  
INFORMATION FOR SEQ ID NO: 16:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 24 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-559-205-16

Query Match 100.0%; Score 10; DB 3; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 18 CTTCTCTTTT 9

RESULT 13  
US-09-417-722-3/c  
Sequence 3, Application US/09417722  
Patent No. 6309837  
GENERAL INFORMATION:  
APPLICANT: Dean, Ralph A.  
APPLICANT: Wang, Yi-Hong  
TITLE OF INVENTION: PCR-based Method for Identifying a Fusarium  
TITLE OF INVENTION: Wilt-Resistant Genotype in Plants  
FILE REFERENCE: PCR ID: Fusarium-resistant genotype  
CURRENT APPLICATION NUMBER: US/09/417,722  
CURRENT FILING DATE: 1999-10-13  
NUMBER OF SEQ ID NOS: 4  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 3  
LENGTH: 24  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence: forward PCR  
OTHER INFORMATION: primer of FM primer pair  
US-09-417-722-3

Query Match 100.0%; Score 10; DB 4; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 23 CTTCTCTTTT 14

RESULT 14  
5336598-16/c  
Patent No. 5336598  
APPLICANT: KOTZIN, BRIAN L.; MARRACK, PHILIPPA; KAPPLER,  
JOHN; CHOI, YOUNGWON  
TITLE OF INVENTION: METHOD FOR DIAGNOSING A SUPERANTIGEN  
CAUSED PATHOLOGICAL CONDITION VIA ASSAY OF T-CELLS  
NUMBER OF SEQUENCES: 25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/437,370  
FILING DATE: 15-NOV-1989  
SEQ ID NO: 16  
LENGTH: 24  
5336598-16

Query Match 100.0%; Score 10; DB 6; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 18 CTTCTCTTTT 9

Query Match 100.0%; Score 10; DB 6; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
| | | | |  
Db 13 CTTCTCTTTT 4

RESULT 17  
US-08-412-376-23/c  
; Sequence 23, Application US/08412376  
; Patent No. 5849900  
; GENERAL INFORMATION:  
; APPLICANT: Moelling, Karin  
; TITLE OF INVENTION: Inhibition Of Viruses By Antisense  
; TITLE OF INVENTION: Oligomers Capable Of Binding To Polypurine-Rich Tract Of Sin  
; TITLE OF INVENTION: Stranded RNA Or RNA-DNA Hybrids  
; NUMBER OF SEQUENCES: 42  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Woodcock Washburn Kurtz  
; ADDRESSEE: Mackiewicz & No. 5849900ris  
; STREET: One Liberty Place - 46th Floor  
; CITY: Philadelphia  
; STATE: PA  
; COUNTRY: USA  
; ZIP: 19103  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE  
; COMPUTER: IBM PS/2  
; OPERATING SYSTEM: PC-DOS  
; SOFTWARE: WORDPERFECT 5.1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/412.376  
; FILING DATE: Herewith  
; CLASSIFICATION: 514  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/954,184  
; FILING DATE: 29-SEP-1992  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Doreen Yanko Trujillo  
; REGISTRATION NUMBER: 35,719  
; REFERENCE/DOCKET NUMBER: APOL-0021  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (215) 568-3100  
; TELEFAX: (215) 568-3439  
; INFORMATION FOR SEQ ID NO: 23:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 27  
; TYPE: Nucleic Acid  
; STRANDEDNESS: Single  
; TOPOLOGY: Linear  
; ANTI-SENSE: Yes

Query Match 100.0%; Score 10; DB 2; Length 27;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
| | | | |  
Db 22 CTTCTCTTTT 13

RESULT 18  
US-08-584-040-7049/c  
; Sequence 7049, Application US/08584040  
; Patent No. 6346398  
; GENERAL INFORMATION:  
; APPLICANT: Pavco, Pamela  
; APPLICANT: McSwiggen, James  
; APPLICANT: Stinchcomb, Dan T.  
; APPLICANT: Escobedo, Jaime

RESULT 15  
US-08-628-417-4  
; Sequence 4, Application US/08628417  
; Patent No. 5627054  
; GENERAL INFORMATION:  
; APPLICANT: GILLESPIE, DAVID  
; TITLE OF INVENTION: COMPETITOR PRIMER ASYMMETRIC  
; TITLE OF INVENTION: POLYMERASE CHAIN REACTION  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: U.S. ARMY CHEMICAL AND BIOLOGICAL  
; ADDRESSEE: DEFENSE COMMAND  
; STREET: OFFICE OF THE CHIEF COUNSEL (AMSCB-GC)  
; CITY: ABERDEEN PROVING GROUND  
; STATE: MARYLAND  
; COUNTRY: USA  
; ZIP: 21010-5423  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/628,417  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: BIFFONI, ULYSSES J  
; REGISTRATION NUMBER: 39,908  
; REFERENCE/DOCKET NUMBER: DAM 398-94  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 410-671-1158  
; TELEFAX: 410-671-2534  
; INFORMATION FOR SEQ ID NO: 4:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 25 bases  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: oligodeoxynucleotide  
; HYPOTHETICAL: NO  
; ANTI-SENSE: YES

Query Match 100.0%; Score 10; DB 1; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
| | | | |  
Db 8 CTTCTCTTTT 17

RESULT 16  
5217891-9/c  
; Patent No. 5217891  
; APPLICANT: BRAKE, ANTHONY J.;VAN DEN BERG, JOHAN A.  
; TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES  
; A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS  
; POLYPEPTIDES  
; NUMBER OF SEQUENCES: 23  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/507,398  
; FILING DATE: 09-APR-1990  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 78,551  
; FILING DATE: 28-JUL-1987  
; SEQ ID NO: 9:  
; LENGTH: 25  
5217891-9

; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: TREATMENT OF DISEASES OR
; TITLE OF INVENTION: CONDITIONS RELATED TO LEVELS
; TITLE OF INVENTION: OF VASCULAR ENDOTHELIAL
; TITLE OF INVENTION: GROWTH FACTOR
; NUMBER OF SEQUENCES: 8502
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071-2066
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: Word Perfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/584,040
; FILING DATE: January 11, 1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/005,974
; FILING DATE: October 26, 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard J.
; REGISTRATION NUMBER: 321,327
; REFERENCE/DOCKET NUMBER: 218/064
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 7049:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; OTHER INFORMATION: The letter "N" represents the stem II region
; OTHER INFORMATION: of an HH ribozyme.
US-08-584-040-7049

Query Match 100.0%; Score 10; DB 4; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 27 CTTCTCTTTT 18

RESULT 19
US-09-270-542-133
; Sequence 133, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, Lawrence
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 133
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-134

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 20
US-09-270-542-132
; Sequence 132, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, Lawrence
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 132
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-133

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 21
US-09-270-542-134
; Sequence 134, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, Lawrence
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 134
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-134

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 8 CTTCTCTTTT 17

## RESULT 22

US-09-270-542-135  
; Sequence 135, Application US/09270542  
; Patent No. 6322976  
; GENERAL INFORMATION:  
; APPLICANT: Altman, Timothy  
; APPLICANT: Scott, James  
; APPLICANT: Stanton, Lawrence  
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and  
; FILE REFERENCE: 4198/78179  
; CURRENT APPLICATION NUMBER: US/09/270,542  
; CURRENT FILING DATE: 1999-03-17  
; EARLIER APPLICATION NUMBER: 09/221,222  
; EARLIER FILING DATE: 1999-12-23  
; NUMBER OF SEQ ID NOS: 207  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 135  
; LENGTH: 30  
; TYPE: DNA  
; ORGANISM: Rattus norvegicus  
US-09-270-542-135

Query Match 100.0%; Score 10; DB 4; Length 30;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 8 CTTCTCTTTT 17

## RESULT 23

US-08-961-083-359/c  
; Sequence 359, Application US/08961083  
; Patent No. 6159469  
; GENERAL INFORMATION:  
; APPLICANT: Choi et. al.  
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines  
; NUMBER OF SEQUENCES: 452  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Human Genome Sciences, Inc.  
; STREET: 9410 Key West Avenue  
; CITY: Rockville  
; STATE: Maryland  
; COUNTRY: USA  
; ZIP: 20850  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage  
; COMPUTER: HP Vectra 486/33  
; OPERATING SYSTEM: MSDOS version 6.2  
; SOFTWARE: ASCII Text  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/961,083  
; FILING DATE:  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER:  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Brookes, A. Anders  
; REGISTRATION NUMBER: 36,373  
; REFERENCE/DOCKET NUMBER: PB340P2  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (301) 309-8504  
; TELEFAX: (301) 309-8512  
; INFORMATION FOR SEQ ID NO: 359:  
; SEQUENCE CHARACTERISTICS:

; LENGTH: 33 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
US-08-961-083-359

Query Match 100.0%; Score 10; DB 3; Length 33;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 23 CTTCTCTTTT 14

## RESULT 24

US-09-536-784-359/c  
; Sequence 359, Application US/09536784  
; Patent No. 6573082  
; GENERAL INFORMATION:  
; APPLICANT: Choi et. al.  
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines  
; NUMBER OF SEQUENCES: 452  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Human Genome Sciences, Inc.  
; STREET: 9410 Key West Avenue  
; CITY: Rockville  
; STATE: Maryland  
; COUNTRY: USA  
; ZIP: 20850  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage  
; COMPUTER: HP Vectra 486/33  
; OPERATING SYSTEM: MSDOS version 6.2  
; SOFTWARE: ASCII Text  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/536,784  
; FILING DATE: 30-Oct-1997  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/961,083  
; FILING DATE: OCT-30-1997  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Michelle S. Marks  
; REGISTRATION NUMBER: 41,971  
; REFERENCE/DOCKET NUMBER: PB340P3  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (301) 309-8504  
; TELEFAX: (301) 309-8512  
; INFORMATION FOR SEQ ID NO: 359:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 33 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:  
US-09-536-784-359

Query Match 100.0%; Score 10; DB 4; Length 33;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 23 CTTCTCTTTT 14

## RESULT 25

US-09-371-772B-10906/c  
; Sequence 10906, Application US/09371772B  
; Patent No. 6566127  
; GENERAL INFORMATION:  
; APPLICANT: Ribozyne Pharmaceuticals, Inc.

APPLICANT: Pavco, Pam  
APPLICANT: McSwiggan, Jim  
APPLICANT: Stinchcomb, Dan  
APPLICANT: Escobedo, Jaime  
TITLE OF INVENTION: Method and Reagent for the Treatment of Diseases or Conditions Related to the Growth of Vascular Endothelial Growth Factor Receptor  
FILE REFERENCE: MBH00,876-J (237/198)  
CURRENT APPLICATION NUMBER: US/09/371,772B  
CURRENT FILING DATE: 1999-08-10  
PRIOR APPLICATION NUMBER: US 60/005,974  
PRIOR FILING DATE: 1995-10-26  
PRIOR APPLICATION NUMBER: US 08/584,040  
PRIOR FILING DATE: 1996-01-08  
NUMBER OF SEQ ID NOS: 14225  
SOFTWARE: PatentIn version 3.0  
SEQ ID NO 10906  
LENGTH: 38  
TYPE: RNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid  
US-09-371-772B-10906

Query Match 100.0%; Score 10; DB 4; Length 38;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 38 CTTCTCTTTT 29

RESULT 26  
5217891-13/c  
Patent No. 5217891  
APPLICANT: BRAKE, ANTHONY J.; VAN DEN BERG, JOHAN A.  
TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES  
A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS  
POLYPEPTIDES  
NUMBER OF SEQUENCES: 23  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/507,398  
FILING DATE: 09-APR-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 78,551  
FILING DATE: 28-JUL-1987  
SEQ ID NO:13:  
LENGTH: 39  
5217891-13

Query Match 100.0%; Score 10; DB 6; Length 39;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 13 CTTCTCTTTT 4

RESULT 27  
US-09-277-016-20  
Sequence 20, Application US/09277016  
Patent No. 6143529  
GENERAL INFORMATION:  
APPLICANT: Lapidus, Stanley N  
APPLICANT: Shuber, Anthony P  
TITLE OF INVENTION: Methods for improving sensitivity and specificity of  
screening assays  
FILE REFERENCE: EXT-030  
CURRENT APPLICATION NUMBER: US/09/277,016  
CURRENT FILING DATE: 1999-03-26  
EARLIER APPLICATION NUMBER: 08/876,857  
EARLIER FILING DATE: 1997-06-16

EARLIER APPLICATION NUMBER: 08/700,583  
EARLIER FILING DATE: 1996-08-14  
NUMBER OF SEQ ID NOS: 37  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 20  
LENGTH: 40  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence:PCR-G-FOR (p53)  
OTHER INFORMATION: Exon 8)  
US-09-277-016-20

Query Match 100.0%; Score 10; DB 3; Length 40;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 31 CTTCTCTTTT 40

RESULT 28  
US-07-908-317-33  
Sequence 33, Application US/07908317  
Patent No. 5420027  
GENERAL INFORMATION:  
APPLICANT: FISHER, CHARLES W.  
APPLICANT: BARNES, HENRY J.  
APPLICANT: ESTABROOK, RONALD W.  
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR  
THE EXPRESSION OF BILOGICALLY  
TITLE OF INVENTION: ACTIVE FUSION PROTEINS COMPRISING A  
TITLE OF INVENTION: EUKARYOTIC CYTOCHROME P450 FUSED TO  
TITLE OF INVENTION: A REDUCTASE IN BACTERIA  
NUMBER OF SEQUENCES: 34  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: ARNOLD, WHITE & DURKEE  
STREET: P.O. BOX 4433  
CITY: HOUSTON  
STATE: TEXAS  
COUNTRY: USA  
ZIP: 77210  
COMPUTER READABLE FORM:  
MEDIUM TYPE: FLOPPY DISK  
COMPUTER: IBM PC COMPATIBLE  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: WORDPERFECT 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/908,317  
FILING DATE: 19920702  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: PARKER, DAVID L.  
REGISTRATION NUMBER: 32,165  
REFERENCE/DOCKET NUMBER: UTSD:292/PAR  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 512-320-7200  
TELEFAX: 512-474-7577  
TELEX: NOT APPLICABLE  
INFORMATION FOR SEQ ID NO: 33:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 42 base pairs  
TYPE: NUCLEIC ACID  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-07-908-317-33

Query Match 100.0%; Score 10; DB 1; Length 42;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

```

RESULT 32
US-09-671-317-765
; Sequence 765, Application US/09671317
; Patent No. 6528260
; GENERAL INFORMATION:
;
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Chugueletet, Lydie
; APPLICANT: Cohen, Annick
; TITLE OF INVENTION: BIALLELIC MARKER
; FILE REFERENCE: 62 US3 CIP
; CURRENT APPLICATION NUMBER: US/09/671
; CURRENT FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US 09/536
; PRIOR FILING DATE: 2000-03-23
; PRIOR APPLICATION NUMBER: PCT/IB00/
; PRIOR FILING DATE: 2000-03-24
; PRIOR APPLICATION NUMBER: US 60/126
; PRIOR FILING DATE: 1999-03-25
; PRIOR APPLICATION NUMBER: US 60/131
; PRIOR FILING DATE: 1999-04-30
; NUMBER OF SEQ ID NOS: 977
; SOFTWARE: Patent.pm
; SEQ ID NO 765
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens

```

```

;
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 12-602-196 : polymorphic base C or T
US-09-171-317-765
; FEATURE:

```

Query Match 100.0%; Score 10; DB 4; Length 47;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 1 CTTCTCTTTT 10

RESULT 33  
US-09-422-978-522/c  
. Sequence 522. Application US/09422978

```

; Patent No. 6537751
;
; GENERAL INFORMATION:
;
; APPLICANT: Cohen, Daniel
;
; APPLICANT: Blumenfeld, Marta
;
; APPLICANT: Chumakov, Ilya
;
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
;
; FILE REFERENCE: GENSET.020CPI
;
; CURRENT APPLICATION NUMBER: US/09/422,978
;
; CURRENT FILING DATE: 1999-10-20
;
; EARLIER APPLICATION NUMBER: US 09/298,850
;
; EARLIER FILING DATE: 1999-04-21
;
; EARLIER APPLICATION NUMBER: US 60/109,732
;
; EARLIER FILING DATE: 1998-11-23
;
; EARLIER APPLICATION NUMBER: US 60/082,614
;
; EARLIER FILING DATE: 1998-04-21
;
; NUMBER OF SEQ ID NOS: 11796
;
; SEQ ID NO 522

```

```

;
; LENGTH: 47
;
; TYPE: DNA
;
; ORGANISM: Homo Sapiens
;
; FEATURE:
;
; NAME/KEY: allele
;
; LOCATION: 24

```

US-09-422-978-522

Query Match 100.0%; Score 10; DB 4; Length 47;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels

Qy	1	CTTCTCTTT	10
Db	15	CTTCTCTTT <td>6</td>	6

RESULT 34  
US-09-422-978-1282  
; Sequence 1282, Application US/09422978  
: Patent NO. 6537751

```

/ GENERAL INFORMATION:
/ APPLICANT: Cohen, Daniel
/ APPLICANT: Blumenfeld, Marta
/ APPLICANT: Chumakov, Il'ya
/ TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
/ FILE REFERENCE: GENSET:020Cp1
/ CURRENT APPLICATION NUMBER: US/09/422,978
/ CURRENT FILING DATE: 1998-10-20
/ EARLIER APPLICATION NUMBER: US 09/298,850
/ EARLIER FILING DATE: 1998-04-21
/ EARLIER APPLICATION NUMBER: US 60/109,732
/ EARLIER FILING DATE: 1998-11-23
/ EARLIER APPLICATION NUMBER: US 60/082,614
/ EARLIER FILING DATE: 1998-04-21
/ NUMBER OF SEQ ID NOS: 11796
/ SEQ ID NO 1282

```

```

1 LENGTH: 47
2 TYPE: DNA
3 ORGANISM: Homo Sapiens
4 FEATURE:
5 NAME/KEY: allele
6 LOCATION: 24
7 OTHER INFORMATION: 99-23604-208 : polymorphic base G or T
US-09-432-978-1282

```

Query Match 100.0%; Score 10; DB 4; Length 47;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels

Qy 1 CTCTCTTTT 10  
db 13 CTCTCTTTT 22

RESULT 35

```

US-09-422-978-1546
; Sequence 1546, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020CPI
; CURRENT APPLICATION NUMBER: US/09/422,978
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1999-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 1546
; LENGTH: 47

```

```

; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-
US-09-422-978-1546

```

Query Match 100.0%; Score 10; DB 4; Length 47;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy	1	CTTCTCTTTT	10
Db	9	CTTCTCTTTT <td>18</td>	18

RESULT 36  
US-09-422-978-2472/c  
; Sequence 2472, Application US/094222978  
: Patent No. 6537751

```

/ GENERAL INFORMATION:
/ APPLICANT: Cohen, Daniel
/ APPLICANT: Blumenfeld, Marta
/ APPLICANT: Chumakov, Ilya
/ TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
/ FILE REFERENCE: GENSET.020CP1
/ CURRENT APPLICATION NUMBER: US 09/422,978
/ CURRENT FILING DATE: 1999-10-20
/ EARLIER APPLICATION NUMBER: US 09/298,850
/ EARLIER FILING DATE: 1999-04-21
/ EARLIER APPLICATION NUMBER: US 60/109,732
/ EARLIER FILING DATE: 1998-11-23
/ EARLIER APPLICATION NUMBER: US 60/082,614

```

Wed Oct 29 15:38:01 2003

```

; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 2472
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; FEATURE:
; LOCATION: 24
; OTHER INFORMATION: 99-11191-86 : polymorphic base A or G
US-09-422-978-2472

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 45 CTTCTCTTTT 36

RESULT 37
US-09-422-978-3284/c
; Sequence 3284, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020CP1
; CURRENT APPLICATION NUMBER: US/09/422,978
; EARLIER FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1999-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 3284
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-2981-53 : polymorphic base T or C
US-09-422-978-3284

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 38
US-09-461-697-225/c
; Sequence 225, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
```

```

; TITLE OF INVENTION: CELL DEATH
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 225
; LENGTH: 48
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-225

Query Match      100.0%; Score 10; DB 3; Length 48;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 20 CTTCTCTTTT 11

RESULT 39
US-08-477-928A-37/c
; Sequence 37, Application US/08477928A
; Patent No. 6207389
; GENERAL INFORMATION:
; APPLICANT: Dosch, Hans M.
; TITLE OF INVENTION: METHODS FOR CONTROLLING T
; TITLE OF INVENTION: LYMPHOCYTE MEDIATED IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 49
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BAKER & BOTTS
; STREET: 1299 Pennsylvania Avenue
; CITY: Washington D.C.
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 20004-2400
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/477,928A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Remenick, James
; REGISTRATION NUMBER: 36902
; REFERENCE/DOCKET NUMBER: 19060-0105
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 639 7700
; TELEFAX: (202) 639 7890
; INFORMATION FOR SEQ ID NO: 37:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-477-928A-37

Query Match      100.0%; Score 10; DB 3; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 42 CTTCTCTTTT 33

RESULT 40
US-08-910-632-40/c
```



```
; Sequence 40, Application US/08910632B
; Patent No. 6077668
; GENERAL INFORMATION:
; APPLICANT: KOOL, ERIC T.
; TITLE OF INVENTION: HIGHLY SENSITIVE MULTIMERIC NUCLEIC ACID PROBES
; FILE REFERENCE: 220.00010130
; CURRENT APPLICATION NUMBER: US/08/910,632B
; CURRENT FILING DATE: 1997-08-13
; EARLIER APPLICATION NUMBER: 08/805,631
; EARLIER FILING DATE: 1997-02-26
; EARLIER APPLICATION NUMBER: 08/393,439
; EARLIER FILING DATE: 1995-02-23
; EARLIER APPLICATION NUMBER: 08/047,860
; EARLIER FILING DATE: 1993-04-15
; NUMBER OF SEQ ID NOS: 83
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 40
; LENGTH: 53
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: DNA 53mer circle
US-08-910-632-40

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 41
US-08-910-632-41
; Sequence 41, Application US/08910632B
; Patent No. 6077668
; GENERAL INFORMATION:
; APPLICANT: KOOL, ERIC T.
; TITLE OF INVENTION: HIGHLY SENSITIVE MULTIMERIC NUCLEIC ACID PROBES
; FILE REFERENCE: 220.00010130
; CURRENT APPLICATION NUMBER: US/08/910,632B
; CURRENT FILING DATE: 1997-08-13
; EARLIER APPLICATION NUMBER: 08/805,631
; EARLIER FILING DATE: 1997-02-26
; EARLIER APPLICATION NUMBER: 08/393,439
; EARLIER FILING DATE: 1995-02-23
; EARLIER APPLICATION NUMBER: 08/047,860
; EARLIER FILING DATE: 1993-04-15
; NUMBER OF SEQ ID NOS: 83
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 41
; LENGTH: 53
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: stem-loop RNA multimer which binds HIV-1 gag RNA
US-08-910-632-41

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.9e+03;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 38 CUUCUCUUU 47

RESULT 42
US-08-805-631A-40/c
; Sequence 40, Application US/08805631A
; Patent No. 6096880
; GENERAL INFORMATION:
; APPLICANT: UNIVERSITY OF ROCHESTER
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
; NUMBER OF SEQUENCES: 72
; CORRESPONDENCE ADDRESS:
; ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
; STREET: 119 No. 6096880th Fourth Street, Suite 201
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55401
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/805,631A
; FILING DATE: 26-FEB-97
; CLASSIFICATION: 536
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/393,439
; FILING DATE: 23-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/047,860
; FILING DATE: 15-APR-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: SANDBERG, VICTORIA A.
; REGISTRATION NUMBER: 41,287
; REFERENCE/DOCKET NUMBER: 220.00010140
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 612-305-1226
; TELEFAX: 612-305-1228
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 53 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
US-08-805-631A-40

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 43
US-08-805-631A-41
; Sequence 41, Application US/08805631A
; Patent No. 6096880
; GENERAL INFORMATION:
; APPLICANT: UNIVERSITY OF ROCHESTER
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
; NUMBER OF SEQUENCES: 72
; CORRESPONDENCE ADDRESS:
; ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
; STREET: 119 No. 6096880th Fourth Street, Suite 201
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55401
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
```

APPLICATION NUMBER: US/08/805,631A  
FILING DATE: 26-FEB-97  
CLASSIFICATION: 536  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/393,439  
FILING DATE: 23-FEB-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/047,860  
FILING DATE: 15-APR-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: SANDBERG, VICTORIA A.  
REGISTRATION NUMBER: 41,287  
REFERENCE/DOCKET NUMBER: 220.00010140  
TELEPHONE: 612-305-1226  
TELEFAX: 612-305-1228  
INFORMATION FOR SEQ ID NO: 41:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 53 base pairs  
TYPE: nucleic acid  
STRADEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: RNA (genomic)  
US-08-805-631A-41

Query Match 100.0%; Score 10; DB 3; Length 53;  
Best Local Similarity 30.0%; Pred. No. 1.9e+03;  
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|:|::|  
DB 38 CUUCUCUUU 47

RESULT 44  
US-09-569-344-40/c  
Sequence 40, Application US/09569344  
Patent No. 6368802  
GENERAL INFORMATION:  
APPLICANT: UNIVERSITY OF ROCHESTER  
TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND  
DNA  
NUMBER OF SEQUENCES: 72  
CORRESPONDENCE ADDRESS:  
ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.  
STREET: 119 No. 6368802th Fourth Street, Suite 201  
CITY: Minneapolis  
STATE: Minnesota  
COUNTRY: USA  
ZIP: 55401  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/569,344  
FILING DATE: 11-May-2000  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/805,631  
FILING DATE: 26-FEB-97  
APPLICATION NUMBER: US 08/393,439  
FILING DATE: 23-FEB-1995  
APPLICATION NUMBER: US 08/047,860  
FILING DATE: 15-APR-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: SANDBERG, VICTORIA A.  
REGISTRATION NUMBER: 41,287  
REFERENCE/DOCKET NUMBER: 220.00010140  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 612-305-1226  
TELEFAX: 612-305-1228

INFORMATION FOR SEQ ID NO: 40:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 53 base pairs  
TYPE: nucleic acid  
STRADEDNESS: single  
TOPOLOGY: circular  
MOLECULE TYPE: DNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 40:  
US-09-569-344-40

Query Match 100.0%; Score 10; DB 4; Length 53;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|:|::|  
DB 25 CTTCTCTTTT 16

RESULT 45  
US-09-569-344-41  
Sequence 41, Application US/09569344  
Patent No. 6368802  
GENERAL INFORMATION:  
APPLICANT: UNIVERSITY OF ROCHESTER  
TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND  
DNA  
NUMBER OF SEQUENCES: 72  
CORRESPONDENCE ADDRESS:  
ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.  
STREET: 119 No. 6368802th Fourth Street, Suite 201  
CITY: Minneapolis  
STATE: Minnesota  
COUNTRY: USA  
ZIP: 55401  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/569,344  
FILING DATE: 11-May-2000  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/805,631  
FILING DATE: 26-FEB-97  
APPLICATION NUMBER: US 08/393,439  
FILING DATE: 23-FEB-1995  
APPLICATION NUMBER: US 08/047,860  
FILING DATE: 15-APR-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: SANDBERG, VICTORIA A.  
REGISTRATION NUMBER: 41,287  
REFERENCE/DOCKET NUMBER: 220.00010140  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 612-305-1226  
TELEFAX: 612-305-1228  
INFORMATION FOR SEQ ID NO: 41:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 53 base pairs  
TYPE: nucleic acid  
STRADEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: RNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 41:  
US-09-569-344-41

Query Match 100.0%; Score 10; DB 4; Length 53;  
Best Local Similarity 30.0%; Pred. No. 1.9e+03;  
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|:|::|  
DB 38 CUUCUCUUU 47

RESULT 44  
US-09-569-344-40/c  
Sequence 40, Application US/09569344  
Patent No. 6368802  
GENERAL INFORMATION:  
APPLICANT: UNIVERSITY OF ROCHESTER  
TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND  
DNA  
NUMBER OF SEQUENCES: 72  
CORRESPONDENCE ADDRESS:  
ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.  
STREET: 119 No. 6368802th Fourth Street, Suite 201  
CITY: Minneapolis  
STATE: Minnesota  
COUNTRY: USA  
ZIP: 55401  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/569,344  
FILING DATE: 11-May-2000  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/805,631  
FILING DATE: 26-FEB-97  
APPLICATION NUMBER: US 08/393,439  
FILING DATE: 23-FEB-1995  
APPLICATION NUMBER: US 08/047,860  
FILING DATE: 15-APR-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: SANDBERG, VICTORIA A.  
REGISTRATION NUMBER: 41,287  
REFERENCE/DOCKET NUMBER: 220.00010140  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 612-305-1226  
TELEFAX: 612-305-1228  
INFORMATION FOR SEQ ID NO: 41:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 53 base pairs  
TYPE: nucleic acid  
STRADEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: RNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 41:  
US-09-569-344-41

Query Match 100.0%; Score 10; DB 4; Length 53;  
Best Local Similarity 30.0%; Pred. No. 1.9e+03;  
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10



```
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/585,684
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2533:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-09-038-073-2533

Query Match 100.0%; Score 10; DB 3; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 49
US-09-038-073-2710/c
; Sequence 2710, Application US/09038073
; Patent No. 6194150
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Daniel T.
; APPLICANT: McSwigen, James
; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: INDUCTION OF GRAFT TOLERANCE
; TITLE OF INVENTION: AND REVERSAL OF IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 2751
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; STREET: Suite 4700
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: FastSeq Version 1.5
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/038,073
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/585,684
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2710:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
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; STRANDEDNESS: single
; TOPOLOGY: linear
US-09-038-073-2710

Query Match 100.0%; Score 10; DB 3; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 50
US-09-446-047A-1
; Sequence 1, Application US/09446047A
; Patent No. 6379924
; GENERAL INFORMATION:
; APPLICANT: Darrell Sleep
; APPLICANT: Delta Biotechnology Limited
; TITLE OF INVENTION: Improved Protein Expression Strains
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Aventis Behring LLC
; STREET: 1020 First Avenue
; CITY: King of Prussia
; STATE: Pennsylvania
; COUNTRY: United States of America
; ZIP: 19406-1310
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/446,047A
; FILING DATE: 15-Dec-1999
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 70 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "PCR primer"
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-446-047A-1

Query Match 100.0%; Score 10; DB 4; Length 70;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 30 CTTCTCTTTT 39

RESULT 51
US-08-117-374-5
; Sequence 5, Application US/08117374
; Patent No. 5362865
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5362865-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5362865th
; CITY: St. Louis
; STATE: Missouri
```

; COUNTRY: USA  
; ZIP: 63198  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA: US/08/117,374  
; FILING DATE: 02-SEP-1993  
; CLASSIFICATION: 800  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Strode, Janelle D.  
; REGISTRATION NUMBER: 34,738  
; REFERENCE/DOCKET NUMBER: 38-21(10531)A  
; TELEPHONE: (314)537-6224  
; TELEFAX: (314)537-6047  
; INFORMATION FOR SEQ ID NO: 5:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 71 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; US-117-374-5  
Query Match 100.0%; Score 10; DB 1; Length 71;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 52

RESULT 52  
US-08-280-263-5  
; Sequence 5, Application US/08280263  
; Patent No. 5659122  
; GENERAL INFORMATION:  
; APPLICANT: Austin, Glenn D.  
; TITLE OF INVENTION: Enhanced Expression in Plants Using  
; NUMBER OF SEQUENCES: 22  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Janelle D. Strode, Monsanto Company, B84F  
; STREET: 700 Chesterfield Parkway No. 5659122th  
; CITY: St. Louis  
; STATE: Missouri  
; COUNTRY: USA  
; ZIP: 63198  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; FILING DATE: 25-JUL-1994  
; CLASSIFICATION: 800  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/117,374  
; FILING DATE: 02-SEP-1993  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Strode, Janelle D.  
; REGISTRATION NUMBER: 34,738  
; REFERENCE/DOCKET NUMBER: 38-21(10531)A  
; TELEPHONE: (314)537-6224  
; TELEFAX: (314)537-6047  
; INFORMATION FOR SEQ ID NO: 5:  
; SEQUENCE CHARACTERISTICS:

; LENGTH: 71 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; US-08-280-263-5

Query Match 100.0%; Score 10; DB 1; Length 71;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 52

RESULT 53  
US-08-597-325-5  
; Sequence 5, Application US/08597325  
; Patent No. 6018100  
; GENERAL INFORMATION:  
; APPLICANT: Rogers, Stephen G.  
; TITLE OF INVENTION: Promoter for Transgenic Plants  
; NUMBER OF SEQUENCES: 11  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Janelle D. Strode, Monsanto Company, B84F  
; STREET: 700 Chesterfield Parkway No. 6018100th  
; CITY: St. Louis  
; STATE: Missouri  
; COUNTRY: USA  
; ZIP: 63198  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA: US/08/597,325  
; FILING DATE: 08/08/597,325  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/366,240  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Strode, Janelle D.  
; REGISTRATION NUMBER: 34,738  
; REFERENCE/DOCKET NUMBER: 38-21(10647)A  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (314)537-6224  
; TELEFAX: (314)537-6047  
; INFORMATION FOR SEQ ID NO: 5:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 71 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; US-08-597-325-5

Query Match 100.0%; Score 10; DB 3; Length 71;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 52

RESULT 54  
US-08-597-325-5  
; Sequence 5, Application US/08597325  
; Patent No. 6018100  
; GENERAL INFORMATION:

APPLICANT: Rogers, Stephen G.  
TITLE OF INVENTION: Promoter for Transgenic Plants  
NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F  
STREET: 700 Chesterfield Parkway No. 6018100th  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63198  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA: 08/597,325  
APPLICATION NUMBER: US/08/597,325  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/366,240  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Strode, Janelle D.  
REGISTRATION NUMBER: 34,738  
REFERENCE/DOCKET NUMBER: 38-21(10647)A  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (314)537-6224  
TELEFAX: (314)537-6047  
INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 71 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-597-325-5

Query Match 100.0%; Score 10; DB 3; Length 71;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 52

RESULT 55  
PCT-US94-10256-5  
Sequence 5, Application PC/TUS9410256  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
TITLE OF INVENTION: Non-translated Leader Sequences  
NUMBER OF SEQUENCES: 22  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25 (EPO)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US94/10256  
FILING DATE: 01-SEPT-1994  
CLASSIFICATION:  
INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 71 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
PCT-US94-10256-5

Query Match 100.0%; Score 10; DB 5; Length 71;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 52

RESULT 56  
US-08-117-374-6/c  
Sequence 6, Application US/08117374  
Patent No. 5362865  
GENERAL INFORMATION:  
APPLICANT: Austin, Glenn D.  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
TITLE OF INVENTION: No. 5362865-translated Leader Sequences  
NUMBER OF SEQUENCES: 22  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F  
STREET: 700 Chesterfield Parkway No. 5362865th  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63198  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/117,374  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Strode, Janelle D.  
REGISTRATION NUMBER: 34,738  
REFERENCE/DOCKET NUMBER: 38-21(10531)A  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (314)537-6224  
TELEFAX: (314)537-6047  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 75 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-117-374-6

Query Match 100.0%; Score 10; DB 1; Length 75;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
Db 33 CTTCTCTTTT 24

RESULT 57  
US-08-280-263-6/c  
Sequence 6, Application US/08280263  
Patent No. 5659122  
GENERAL INFORMATION:  
APPLICANT: Austin, Glenn D.  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
TITLE OF INVENTION: No. 5659122-translated Leader Sequences  
NUMBER OF SEQUENCES: 22  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F  
STREET: 700 Chesterfield Parkway No. 5659122th  
CITY: St. Louis  
STATE: Missouri

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; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/280,263
; FILING DATE: 25-JUL-1994
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/117,374
; FILING DATE: 02-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
```

```
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 75 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-597-325-6

Query Match 100.0%; Score 10; DB 3; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 33 CTTCTCTTTT 24

RESULT 59
US-08-597-325-6/c
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
```

```
RESULT 60
PCT-US94-10256-6/c
; Sequence 6, Application PC/TUS9410256
```

GENERAL INFORMATION:  
APPLICANT: Enhanced Expression in Plants Using  
TITLE OF INVENTION: Non-translated Leader Sequences  
NUMBER OF SEQUENCES: 22  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25 (BPO)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US94/10256  
FILING DATE: 01-SEPT-1994  
CLASSIFICATION:  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 75 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
PCT-US94-10256-6

Query Match 100.0%; Score 10; DB 5; Length 75;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 33 CTTCTCTTTT 24

RESULT 61  
US-08-117-374-19  
Sequence 19, Application US/08117374  
Patent No. 5362865  
GENERAL INFORMATION:  
APPLICANT: Austin, Glenn D.  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
NUMBER OF SEQUENCES: 22  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F  
STREET: 700 Chesterfield Parkway No. 5362865th  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63198  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/117,374  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Strode, Janelle D.  
REGISTRATION NUMBER: 34,738  
REFERENCE/DOCKET NUMBER: 38-21(10531)A  
TELEPHONE: (314)537-6224  
TELEFAX: (314)537-6047  
INFORMATION FOR SEQ ID NO: 19:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 78 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-117-374-19

Query Match 100.0%; Score 10; DB 1; Length 78;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
|||||  
Db 43 CTTCTCTTTT 52  
RESULT 62  
US-08-280-263-19  
Sequence 19, Application US/08280263  
Patent No. 5659122  
GENERAL INFORMATION:  
APPLICANT: Austin, Glenn D.  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
NUMBER OF SEQUENCES: 22  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F  
STREET: 700 Chesterfield Parkway No. 5659122th  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63198  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/280,263  
FILING DATE: 25-JUL-1994  
CLASSIFICATION: 800  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/117,374  
FILING DATE: 02-SEP-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Strode, Janelle D.  
REGISTRATION NUMBER: 34,738  
REFERENCE/DOCKET NUMBER: 38-21(10531)A  
TELEPHONE: (314)537-6224  
TELEFAX: (314)537-6047  
INFORMATION FOR SEQ ID NO: 19:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 78 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-280-263-19

Query Match 100.0%; Score 10; DB 1; Length 78;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||  
Db 43 CTTCTCTTTT 52

RESULT 63  
PCT-US94-10256-19  
Sequence 19, Application PC/TUS9410256  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: Enhanced Expression in Plants Using  
NUMBER OF SEQUENCES: 22  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
US-08-280-263-19



```

; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/10256
; FILING DATE: 01-SEPT-1994
; CLASSIFICATION:
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 78 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
PCT-US94-10256-19

```

```

Query Match 100.0%; Score 10; DB 5; Length 78;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 CTTCTCTTTT 10
Db 43 CTTCTCTTTT 52

```

```

RESULT 64
US-08-117-374-20/c
; Sequence 20, Application US/08117374
; Patent No. 5362865
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5362865-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESS: Janelle D. Strobe, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5362865th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/117,374
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Strobe, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 20:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 82 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-117-374-20

```

```

Query Match 100.0%; Score 10; DB 1; Length 82;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 CTTCTCTTTT 10
Db 40 CTTCTCTTTT 31

```

```

RESULT 65
US-08-280-263-20/c
; Sequence 20, Application US/08280263
; Patent No. 5659122
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5659122-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESS: Janelle D. Strobe, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5659122th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/280,263
; FILING DATE: 25-JUL-1994
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/117,374
; FILING DATE: 02-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Strobe, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 20:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 82 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-280-263-20

```

```

Query Match 100.0%; Score 10; DB 1; Length 82;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 CTTCTCTTTT 10
Db 40 CTTCTCTTTT 31

```

```

RESULT 66
PCT-US94-10256-20/c
; Sequence 20, Application PC/TUS9410256
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/10256
; FILING DATE: 01-SEPT-1994
; CLASSIFICATION:
; INFORMATION FOR SEQ ID NO: 20:
; SEQUENCE CHARACTERISTICS:

```

```
; LENGTH: 82 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
PCT-US94-10256-20

Query Match          100.0%; Score 10; DB 5; Length 82;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 40 CTTCTCTTTT 31

RESULT 67
US-09-174-465D-11/c
; Sequence 11, Application US/09174465D
; Patent No. 6180364
; GENERAL INFORMATION:
; APPLICANT: KOMAN, Ahment
; APPLICANT: CHASSIN, Dorine
; APPLICANT: BELLET, Dominique
; TITLE OF INVENTION: NEW PROTEIN CALLED EPIL/PLACENTIN, PROCESS FOR THE
; TITLE OF INVENTION: PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING SUCH, DNA CODING FOR SAID
; TITLE OF INVENTION: PROTEIN
; FILE REFERENCE: 017753-103
; CURRENT APPLICATION NUMBER: US/09/174,465D
; CURRENT FILING DATE: 1998-10-19
; PRIOR APPLICATION NUMBER: US 08/482,842
; PRIOR FILING DATE: 1995-06-07
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 11
; LENGTH: 93
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(93)
; OTHER INFORMATION: Description of Unknown Organism:EPIL - Early
; OTHER INFORMATION: Placenta Insulin-Like peptide
US-09-174-465D-11

Query Match          100.0%; Score 10; DB 3; Length 93;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 17 CTTCTCTTTT 8

RESULT 68
US-09-599-564A-11/c
; Sequence 11, Application US/09599564A
; Patent No. 6362318
; GENERAL INFORMATION:
; APPLICANT: KOMAN, Ahment
; APPLICANT: CHASSIN, Dorine
; APPLICANT: BELLET, Dominique
; TITLE OF INVENTION: NEW PROTEIN CALLED EPIL/PLACENTIN, PROCESS FOR THE
; TITLE OF INVENTION: PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING SUCH, DNA CODING FOR SAID
; TITLE OF INVENTION: PROTEIN
; FILE REFERENCE: 017753-127
; CURRENT APPLICATION NUMBER: US/09/599,564A
; CURRENT FILING DATE: 2000-06-23
; PRIOR APPLICATION NUMBER: 09/174,465
; PRIOR FILING DATE: 1998-10-19
; PRIOR APPLICATION NUMBER: US 08/482,842

Query Match          100.0%; Score 10; DB 3; Length 93;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 17 CTTCTCTTTT 8

RESULT 69
US-09-461-697-223/c
; Sequence 223, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; TITLE OF INVENTION: CELL DEATH
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 223
; LENGTH: 96
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-223

Query Match          100.0%; Score 10; DB 3; Length 96;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 68 CTTCTCTTTT 59

RESULT 70
US-09-144-428-55/c
; Sequence 55, Application US/09144428
; Patent No. 6583108
; GENERAL INFORMATION:
; APPLICANT: BAYER CORPORATION, The
; APPLICANT: TAMBURINI, Paul P
; APPLICANT: DAVIS, Gary
; APPLICANT: DELARIA, Katherine A
; APPLICANT: MARLOR, Christopher W
; APPLICANT: MULLER, Daniel K
; TITLE OF INVENTION: HUMAN BIKUNIN
; NUMBER OF SEQUENCES: 71
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff
```

STREET: 300 S. Wacker Drive Suite 3200  
CITY: CHICAGO  
STATE: ILLINOIS  
COUNTRY: USA  
ZIP: 60606  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/144.428  
FILING DATE: 10-MAR-1997  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: WO PCT/US97/03894  
FILING DATE: 10-MAR-1997  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 60/013.106  
FILING DATE: 11-MAR-1996  
APPLICATION DATA:  
APPLICATION NUMBER: US 60/019.793  
FILING DATE: 14-JUN-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/725.251  
FILING DATE: 04-OCT-1996  
ATTORNEY/AGENT INFORMATION:  
NAME: CHAO, Mark  
REGISTRATION NUMBER: 37,293  
REFERENCE/DOCKET NUMBER: 96,223-II  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 913-0001  
TELEFAX: (312) 913-0002  
INFORMATION FOR SEQ ID NO: 55:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 102 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
US-09-144-428-55

Query Match 100.0%; Score 10; DB 4; Length 102;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 28 CTTCTCTTTT 19

RESULT 71  
US-08-932-082-10  
Sequence 10, Application US/08932082  
Patent No. 6251856  
GENERAL INFORMATION:

APPLICANT: Markussen, Jan  
APPLICANT: Jonassen, Ib  
APPLICANT: Havelund, Svend  
APPLICANT: Brandt, Jakob  
APPLICANT: Kurtzhals, Peter  
APPLICANT: Hansen, Hertz Per  
APPLICANT: Kaarsholm, Niels Christian  
TITLE OF INVENTION: INSULIN DERIVATIVES  
NUMBER OF SEQUENCES: 26  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: No. 6251856 No. 6251856disk of No. 6251856th America, Inc.  
STREET: 405 Lexington Avenue, 64th Floor  
CITY: New York  
STATE: New York  
COUNTRY: United States of America  
ZIP: 10174-6401

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/932,082  
FILING DATE: 12-AUG-1997  
CLASSIFICATION: 514  
ATTORNEY/AGENT INFORMATION:  
NAME: Lambiris, Elias J.  
REGISTRATION NUMBER: 33,728  
REFERENCE/DOCKET NUMBER: 4341.204-US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 212-867-0123  
TELEFAX: 212-878-9655  
INFORMATION FOR SEQ ID NO: 10:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 112 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-08-932-082-10

Query Match 100.0%; Score 10; DB 3; Length 112;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 88 CTTCTCTTTT 97

## RESULT 72

US-09-461-697-221/c  
Sequence 221, Application US/09461697  
Patent No. 6277974

GENERAL INFORMATION:  
APPLICANT: COGENT NEUROSCIENCE, Inc.  
APPLICANT: Lo, Donald C.  
APPLICANT: Barney, Shawn  
APPLICANT: Thomas, Mary Beth  
APPLICANT: Portbury, Stuart D.  
APPLICANT: Purnam, Kasturi  
APPLICANT: Katz, Lawrence C.  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
FILE REFERENCE: 10001-085-999  
CURRENT APPLICATION NUMBER: US/09/461,697  
CURRENT FILING DATE: 1999-12-14  
NUMBER OF SEQ ID NOS: 466  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 221  
LENGTH: 117  
TYPE: DNA  
ORGANISM: Homo sapiens  
US-09-461-697-221

Query Match 100.0%; Score 10; DB 3; Length 117;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 89 CTTCTCTTTT 80

## RESULT 73

US-09-461-697-219/c  
Sequence 219, Application US/09461697  
Patent No. 6277974  
GENERAL INFORMATION:

```
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 219
; LENGTH: 126
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-219

Query Match      100.0%; Score 10; DB 3; Length 126;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 98 CTTCTCTTTT 89

RESULT 74
US-09-441-416A-22/c
; Sequence 22, Application US/09441416A
; Patent No. 6294518
; GENERAL INFORMATION:
; APPLICANT: Potter, David A.
; APPLICANT: Skolnik, Paul R.
; TITLE OF INVENTION: CELL-PERMEABLE PROTEIN INHIBITORS OF
; TITLE OF INVENTION: CALPAIN
; FILE REFERENCE: 00398-140001
; CURRENT APPLICATION NUMBER: US/09/441,416A
; CURRENT FILING DATE: 1999-11-16
; PRIOR APPLICATION NUMBER: US 08/964,302
; PRIOR FILING DATE: 1997-11-04
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22
; LENGTH: 132
; TYPE: DNA
; ORGANISM: Eukaryote
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (7)...(129)
US-09-441-416A-22

Query Match      100.0%; Score 10; DB 3; Length 132;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 79 CTTCTCTTTT 70

RESULT 75
US-08-477-928A-36/c
; Sequence 36, Application US/08477928A
; Patent No. 6207389
; GENERAL INFORMATION:
; APPLICANT: Dosch, Hans M.
; TITLE OF INVENTION: METHODS FOR CONTROLLING T
; TITLE OF INVENTION: LYMPHOCYTE MEDIATED IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 49
```

```
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BAKER & BOTTS
; STREET: 1299 Pennsylvania Avenue
; CITY: Washington D.C.
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 20004-2400
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/477,928A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Remenick, James
; REGISTRATION NUMBER: 36902
; REFERENCE/DOCKET NUMBER: 19060-0105
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 639 7700
; TELEFAX: (202) 639 7890
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-477-928A-36

Query Match      100.0%; Score 10; DB 3; Length 146;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 138 CTTCTCTTTT 129

RESULT 76
US-09-461-697-217/c
; Sequence 217, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 217
; LENGTH: 156
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-217

Query Match      100.0%; Score 10; DB 3; Length 156;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 138 CTTCTCTTTT 129
```

Db 128 CTTCTCTTTT 119

## RESULT 77

US-09-461-697-215/c  
; Sequence 215, Application US/09461697  
; Patent No. 6277974  
; GENERAL INFORMATION:  
; APPLICANT: COGENT NEUROSCIENCE, Inc.  
; APPLICANT: Lo, Donald C.  
; APPLICANT: Barney, Shawn  
; APPLICANT: Thomas, Mary Beth  
; APPLICANT: Portbury, Stuart D.  
; APPLICANT: Purnam, Kasturi  
; APPLICANT: Katz, Lawrence C.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
; FILE REFERENCE: 10001-005-999  
; CURRENT APPLICATION NUMBER: US/09/461,697  
; CURRENT FILING DATE: 1999-12-14  
; NUMBER OF SEQ ID NOS: 466  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 215  
; LENGTH: 174  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-461-697-215

Query Match 100.0%; Score 10; DB 3; Length 174;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 146 CTTCTCTTTT 137

## RESULT 78

US-09-461-697-213/c  
; Sequence 213, Application US/09461697  
; Patent No. 6277974  
; GENERAL INFORMATION:  
; APPLICANT: COGENT NEUROSCIENCE, Inc.  
; APPLICANT: Lo, Donald C.  
; APPLICANT: Barney, Shawn  
; APPLICANT: Thomas, Mary Beth  
; APPLICANT: Portbury, Stuart D.  
; APPLICANT: Purnam, Kasturi  
; APPLICANT: Katz, Lawrence C.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
; FILE REFERENCE: 10001-005-999  
; CURRENT APPLICATION NUMBER: US/09/461,697  
; CURRENT FILING DATE: 1999-12-14  
; NUMBER OF SEQ ID NOS: 466  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 213  
; LENGTH: 189  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-461-697-213

Query Match 100.0%; Score 10; DB 3; Length 189;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 161 CTTCTCTTTT 152

## RESULT 79

US-09-461-697-211/c  
; Sequence 211, Application US/09461697  
; Patent No. 6277974  
; GENERAL INFORMATION:  
; APPLICANT: COGENT NEUROSCIENCE, Inc.  
; APPLICANT: Lo, Donald C.  
; APPLICANT: Barney, Shawn  
; APPLICANT: Thomas, Mary Beth  
; APPLICANT: Portbury, Stuart D.  
; APPLICANT: Purnam, Kasturi  
; APPLICANT: Katz, Lawrence C.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
; FILE REFERENCE: 10001-005-999  
; CURRENT APPLICATION NUMBER: US/09/461,697  
; CURRENT FILING DATE: 1999-12-14  
; NUMBER OF SEQ ID NOS: 466  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 211  
; LENGTH: 195  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-461-697-211

Query Match 100.0%; Score 10; DB 3; Length 195;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 167 CTTCTCTTTT 158

## RESULT 80

US-09-107-532A-2918/c  
; Sequence 2918, Application US/09107532A  
; Patent No. 6583275  
; GENERAL INFORMATION:  
; APPLICANT: Lynn A Doucette-Stamm and David Bush  
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS  
; NUMBER OF SEQUENCES: 7310  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION  
; STREET: 100 Beaver Street  
; CITY: Walcham  
; STATE: Massachusetts  
; COUNTRY: USA  
; ZIP: 02354  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: CD-ROM ISO9660  
; COMPUTER: PC  
; OPERATING SYSTEM: <Unknown>  
; SOFTWARE: ASCII  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/107,532A  
; FILING DATE: 30-Jun-1998  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 60/085,598  
; FILING DATE: 14 May 1998  
; APPLICATION NUMBER: 60/051571  
; FILING DATE: July 2, 1997  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Ariniello, Pamela Deneke  
; REGISTRATION NUMBER: 40,489  
; REFERENCE/DOCKET NUMBER: GTC-012  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (781)893-5007  
; TELEFAX: (781)893-8277  
; INFORMATION FOR SEQ ID NO: 2918:  
; SEQUENCE CHARACTERISTICS:

```
;
; LENGTH: 198 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...198
; SEQUENCE DESCRIPTION: SEQ ID NO: 2918:
US-09-107-532A-2918

Query Match      100.0%; Score 10; DB 4; Length 198;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      126 CTTCTCTTTT 117

RESULT 81
US-09-107-532A-439
; Sequence 439, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 439:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 201 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (B) LOCATION 1...201
; SEQUENCE DESCRIPTION: SEQ ID NO: 2918:
US-09-107-532A-2918

Query Match      100.0%; Score 10; DB 4; Length 198;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      126 CTTCTCTTTT 117

RESULT 82
US-09-107-532A-3253/c
; Sequence 3253, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 3253:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 201 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...201
; SEQUENCE DESCRIPTION: SEQ ID NO: 3253:
US-09-107-532A-3253

Query Match      100.0%; Score 10; DB 4; Length 201;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      178 CTTCTCTTTT 169
```

RESULT 83  
US-09-313-294A-7004  
; Sequence 7004, Application US/09313294A  
; Patent No. 6476212  
; GENERAL INFORMATION:  
; APPLICANT: Lalgudi, Raghunath V.  
; APPLICANT: Ito, Laura Y.  
; APPLICANT: Sherman, Bradley K.  
; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR  
; FILE REFERENCE: PL-0017 US  
; CURRENT APPLICATION NUMBER: US/09/313,294A  
; CURRENT FILING DATE: 1999-05-14  
; NUMBER OF SEQ ID NOS: 7600  
; SOFTWARE: PERL Program  
; SEQ ID NO 7004  
; LENGTH: 206  
; TYPE: DNA  
; ORGANISM: Zea mays  
; FEATURE:  
; NAME/KEY: misc feature  
; OTHER INFORMATION: Incyte ID No. 6476212 700380982H1  
; NAME/KEY: unsure  
; LOCATION: 190, 193  
; OTHER INFORMATION: a, t, c, g, or other  
US-09-313-294A-7004

Query Match 100.0%; Score 10; DB 4; Length 206;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
DB 129 CTTCTCTTTT 138

RESULT 84  
US-09-461-697-209/c  
; Sequence 209, Application US/09461697  
; Patent No. 6277974  
; GENERAL INFORMATION:  
; APPLICANT: COGENT NEUROSCIENCE, Inc.  
; APPLICANT: Lo, Donald C.  
; APPLICANT: Barney, Shawn  
; APPLICANT: Thomas, Mary Beth  
; APPLICANT: Portbury, Stuart D.  
; APPLICANT: Puranam, Kasturi  
; APPLICANT: Katz, Lawrence C.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
; FILE REFERENCE: 10001-005-999  
; CURRENT APPLICATION NUMBER: US/09/461,697  
; CURRENT FILING DATE: 1999-12-14  
; NUMBER OF SEQ ID NOS: 466  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 209  
; LENGTH: 213  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-461-697-209

Query Match 100.0%; Score 10; DB 3; Length 213;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
DB 185 CTTCTCTTTT 176

RESULT 85

5217891-19/c  
; Patent No. 5217891  
; APPLICANT: BRAKE, ANTHONY J.; VAN DEN BERG, JOHAN A.  
; TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES  
; A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS  
; POLYPEPTIDES  
; NUMBER OF SEQUENCES: 23  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/507,398  
; FILING DATE: 09-APR-1990  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 78,551  
; FILING DATE: 28-JUL-1987  
; SEQ ID NO:19:  
; LENGTH: 222  
5217891-19  
Query Match 100.0%; Score 10; DB 6; Length 222;  
Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTTCTCTTTT 10  
DB 37 CTTCTCTTTT 28  
RESULT 86  
US-09-016-434-99/c  
; Sequence 99, Application US/09016434  
; Patent No. 6500938  
; GENERAL INFORMATION:  
; APPLICANT: Janice Au-Young  
; APPLICANT: Jeffrey J. Seilhamer  
; TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF SIGNALING  
; TITLE OF INVENTION: PATHWAY GENE EXPRESSION  
; NUMBER OF SEQUENCES: 1490  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.  
; STREET: 3174 PORTER DRIVE  
; CITY: PALO ALTO  
; STATE: CALIFORNIA  
; COUNTRY: USA  
; ZIP: 94304  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/016,434  
; FILING DATE: HEREMITH  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER:  
; FILING DATE:  
; CLASSIFICATION:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Zeller, Karen J.  
; REGISTRATION NUMBER: 37,071  
; REFERENCE/DOCKET NUMBER: PA-0002 US  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (650) 855-0555  
; TELEFAX: (650) 845-4166  
; INFORMATION FOR SEQ ID NO: 99:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 228 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; IMMEDIATE SOURCE:  
; LIBRARY: LUNGFET03  
; CLONE: 1251228  
US-09-016-434-99

```
Query Match      100.0%; Score 10; DB 4; Length 228;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTTTT 10
Db      58 CTCTCTTTT 49

RESULT 87
US-09-461-697-207/c
; Sequence 207, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 207
; LENGTH: 231
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-207

Query Match      100.0%; Score 10; DB 3; Length 231;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTTTT 10
Db      203 CTCTCTTTT 194

RESULT 88
US-09-107-532A-2834/c
; Sequence 2834, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
```

```
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 2834:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 231 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...231
; SEQUENCE DESCRIPTION: SEQ ID NO: 2834:
US-09-107-532A-2834
; Query Match      100.0%; Score 10; DB 4; Length 231;
; Best Local Similarity 100.0%; Pred. No. 1.8e+03;
; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTTTT 10
Db      79 CTCTCTTTT 70

RESULT 89
US-09-107-532A-724/c
; Sequence 724, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 724:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 234 base pairs
```



```
;
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
;
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (8) LOCATION 1...234
; SEQUENCE DESCRIPTION: SEQ ID NO: 724:
US-09-107-532A-724

Query Match 100.0%; Score 10; DB 4; Length 234;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 118 CTTCTCTTTT 109

RESULT 90
US-09-397-787-56
; Sequence 56, Application US/09397787
; Patent No. 6468758
; GENERAL INFORMATION:
; APPLICANT: Benson, Darin R.
; APPLICANT: Lodes, Michael J.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: King, Gordon E.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
; TITLE OF INVENTION: CANCER THERAPY AND DIAGNOSIS
; FILE REFERENCE: 210121.466C2
; CURRENT APPLICATION NUMBER: US/09/397,787
; CURRENT FILING DATE: 1999-09-16
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 56
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapien
US-09-397-787-56

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 71 CTTCTCTTTT 80

RESULT 91
US-09-389-681-360
; Sequence 360, Application US/09389681A
; Patent No. 6518237
; GENERAL INFORMATION:
; APPLICANT: Yuqui, Jiang
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; TITLE OF INVENTION: COMPOSITIONS OF BREAST CANCER AND METHODS FOR THEIR USE
; FILE REFERENCE: 210121.470C3
; CURRENT APPLICATION NUMBER: US/09/389,681A
; CURRENT FILING DATE: 1999-09-02
; NUMBER OF SEQ ID NOS: 463
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA

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; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-389-681-360

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 113 CTTCTCTTTT 122

RESULT 92
US-09-620-405B-360
; Sequence 360, Application US/09620405B
; Patent No. 6528054
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yuqui
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Hepler, William T.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS OF BREAST CANCER
; FILE REFERENCE: 210121.470C8
; CURRENT APPLICATION NUMBER: US/09/620,405B
; CURRENT FILING DATE: 2000-07-20
; NUMBER OF SEQ ID NOS: 495
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-620-405B-360

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 113 CTTCTCTTTT 122

RESULT 93
US-09-433-826B-360
; Sequence 360, Application US/09433826B
; Patent No. 6579973
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yuqui
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; TITLE OF INVENTION: COMPOSITIONS FOR THE TREATMENT AND
; TITLE OF INVENTION: DIAGNOSIS OF BREAST CANCER AND METHODS FOR THEIR USE
; FILE REFERENCE: 210121.470C4
; CURRENT APPLICATION NUMBER: US/09/433,826B
; CURRENT FILING DATE: 1999-11-03
; NUMBER OF SEQ ID NOS: 474
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
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; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-433-826B-360

Query Match      100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db     113 CTTCTCTTTT 122

RESULT 94
US-09-604-287A-360
; Sequence 360, Application US/09604287A
; Patent No. 6586572
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yugu
; APPLICANT: Dillon, Jenni C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Hepler, William T.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS OF BREAST CANCER
; FILE REFERENCE: 210121.470C7
; CURRENT APPLICATION NUMBER: US/09/604,287A
; CURRENT FILING DATE: 2000-06-22
; NUMBER OF SEQ ID NOS: 489
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-604-287A-360

Query Match      100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db     113 CTTCTCTTTT 122

RESULT 95
US-09-016-434-737/c
; Sequence 737, Application US/09016434
; Patent No. 6500938
; GENERAL INFORMATION:
; APPLICANT: Janice Au-Young
; APPLICANT: Jeffrey J. Seilhamer
; TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF SIGNALING
; TITLE OF INVENTION: PATHWAY GENE EXPRESSION
; NUMBER OF SEQUENCES: 1490
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 PORTER DRIVE
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/098,789
; FILING DATE: Herewith
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: CERRONE, MICHAEL C.
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0547 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166

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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/016,434
; FILING DATE: HEREWITH
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Zeller, Karen J.
; REGISTRATION NUMBER: 37,071
; REFERENCE/DOCKET NUMBER: PA-0002 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166
; INFORMATION FOR SEQ ID NO: 737:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: TMLR2DT01
; CLONE: 395476
US-09-016-434-737

Query Match      100.0%; Score 10; DB 4; Length 247;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db     118 CTTCTCTTTT 109

RESULT 96
US-09-098-789-9
; Sequence 9, Application US/09098789
; Patent No. 6180342
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; APPLICANT: Tang, Y. Tom
; APPLICANT: Lal, Preeti
; APPLICANT: Corley, Neil C.
; APPLICANT: Guegler, Karl J.
; APPLICANT: Patterson, Chandra
; TITLE OF INVENTION: VACULAR PROTON ATPASE SUBUNITS
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 PORTER DRIVE
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/098,789
; FILING DATE: Herewith
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: CERRONE, MICHAEL C.
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0547 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166

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/ INFORMATION FOR SEQ ID NO: 9:  
/ SEQUENCE CHARACTERISTICS:  
/ LENGTH: 251 base pairs  
/ TYPE: nucleic acid  
/ STRANDEDNESS: single  
/ TOPOLOGY: linear  
/ IMMEDIATE SOURCE:  
/ LIBRARY: HIPONON02  
/ CLONE: 2246348CT1  
US-09-098-789-9

Query Match 100.0%; Score 10; DB 3; Length 251;  
Best Local Similarity 100.0%; Pred. No. 1.7e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 108 CTTCTCTTTT 117

RESULT 97  
US-09-702-705-1077/c  
/ Sequence 1077, Application US/09702705  
/ Patent No. 6504010  
/ GENERAL INFORMATION:  
/ APPLICANT: Wang, Tongtong  
/ APPLICANT: Bangur, Chaitanya S.  
/ APPLICANT: Lodes, Michael A.  
/ APPLICANT: Fanger, Gary  
/ APPLICANT: Vedvick, Tom  
/ APPLICANT: Carter, Darrick  
/ APPLICANT: Retter, Marc  
/ APPLICANT: Mannion, Jane  
/ APPLICANT: Fan, Liqun  
/ TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND  
/ FILE REFERENCE: 210121.478C14  
/ CURRENT APPLICATION NUMBER: US/09/702,705  
/ CURRENT FILING DATE: 2000-10-30  
/ NUMBER OF SEQ ID NOS: 1833  
/ SOFTWARE: FastSeq for Windows Version 3.0  
/ SEQ ID NO 1077  
/ LENGTH: 256  
/ TYPE: DNA  
/ ORGANISM: Homo sapien  
US-09-702-705-1077

Query Match 100.0%; Score 10; DB 4; Length 256;  
Best Local Similarity 100.0%; Pred. No. 1.7e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 132 CTTCTCTTTT 123

RESULT 98  
US-09-736-457-1077/c  
/ Sequence 1077, Application US/09736457  
/ Patent No. 6509448  
/ GENERAL INFORMATION:  
/ APPLICANT: Wang, Tongtong  
/ APPLICANT: Bangur, Chaitanya S.  
/ APPLICANT: Lodes, Michael A.  
/ APPLICANT: Fanger, Gary  
/ APPLICANT: Vedvick, Tom  
/ APPLICANT: Carter, Darrick  
/ APPLICANT: Retter, Marc  
/ APPLICANT: Mannion, Jane  
/ APPLICANT: Fan, Liqun  
/ APPLICANT: Wang, Aijun  
/ TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND  
/ FILE REFERENCE: 210121.478C14  
/ CURRENT APPLICATION NUMBER: US/09/702,705  
/ CURRENT FILING DATE: 2000-10-30  
/ NUMBER OF SEQ ID NOS: 1833  
/ SOFTWARE: FastSeq for Windows Version 3.0  
/ SEQ ID NO 1077  
/ LENGTH: 256  
/ TYPE: DNA  
/ ORGANISM: Homo sapien  
US-09-702-705-1077

/ FILE REFERENCE: 210121.478C15  
/ CURRENT APPLICATION NUMBER: US/09/736,457  
/ CURRENT FILING DATE: 2000-12-13  
/ NUMBER OF SEQ ID NOS: 1864  
/ SOFTWARE: FastSeq for Windows Version 3.0  
/ SEQ ID NO 1077  
/ LENGTH: 256  
/ TYPE: DNA  
/ ORGANISM: Homo sapien  
US-09-736-457-1077

Query Match 100.0%; Score 10; DB 4; Length 256;  
Best Local Similarity 100.0%; Pred. No. 1.7e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 132 CTTCTCTTTT 123

RESULT 99  
US-09-313-294A-3143/c  
/ Sequence 3143, Application US/09313294A  
/ Patent No. 6476212  
/ GENERAL INFORMATION:  
/ APPLICANT: Lalgudi, Raghunath V.  
/ APPLICANT: Ito, Laura Y.  
/ APPLICANT: Sherman, Bradley K.  
/ TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR  
/ FILE REFERENCE: PL-0017 US  
/ CURRENT APPLICATION NUMBER: US/09/313,294A  
/ CURRENT FILING DATE: 1999-05-14  
/ NUMBER OF SEQ ID NOS: 7600  
/ SOFTWARE: PERL Program  
/ SEQ ID NO 3143  
/ LENGTH: 263  
/ TYPE: DNA  
/ ORGANISM: Zea mays  
/ NAME/KEY: misc feature  
/ OTHER INFORMATION: Incyte ID No. 6476212 700610958H1  
/ NAME/KEY: unsure  
/ LOCATION: 216, 246  
/ OTHER INFORMATION: a, t, c, g, or other  
US-09-313-294A-3143

Query Match 100.0%; Score 10; DB 4; Length 263;  
Best Local Similarity 100.0%; Pred. No. 1.7e+03;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
Db 166 CTTCTCTTTT 157

RESULT 100  
US-09-134-001C-18/c  
/ Sequence 18, Application US/09134001C  
/ Patent No. 6380370  
/ GENERAL INFORMATION:  
/ APPLICANT: Lynn Doucette-Stamm et al  
/ TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO STAPHYLOCOCCUS  
/ FILE REFERENCE: GTC-007  
/ CURRENT APPLICATION NUMBER: US/09/134,001C  
/ CURRENT FILING DATE: 1998-08-13  
/ PRIOR APPLICATION NUMBER: US 60/064,964  
/ PRIOR FILING DATE: 1997-11-08  
/ PRIOR APPLICATION NUMBER: US 60/055,779  
/ PRIOR FILING DATE: 1997-08-14  
/ NUMBER OF SEQ ID NOS: 5674  
/ SEQ ID NO 18  
/ LENGTH: 267

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; TYPE: DNA
; ORGANISM: Staphylococcus epidermidis
US-09-134-001C-18

Query Match      100.0%; Score 10; DB 4; Length 267;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      159 CTTCTCTTTT 150
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Job time : 96 secs
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GenCore version 5.1.6  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 17:14:13 ; Search time 1747 Seconds  
(without alignments)  
15.351 Million cell updates/sec.

Title: US-09-335-032-71

Perfect score: 10

Sequence: 1 cttctctttt 10

Scoring table: OLIGO\_NUC

Gapop 60.0 , Gapext 60.0

Searched: 1792395 seqs, 1340900451 residues

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

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C 6	10	100.0	17	10	US-09-827-998-292
C 7	10	100.0	17	10	US-09-827-998-293
C 8	10	100.0	17	10	US-09-827-998-294
C 9	10	100.0	17	11	US-09-930-423-1740
C 10	10	100.0	17	11	US-09-930-423-1741
C 11	10	100.0	17	12	US-09-745-237A-1740
C 12	10	100.0	17	12	US-09-745-237A-1741
C 13	10	100.0	17	14	US-09-745-237A-1741
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14	US-10-060-756A-1725	Sequence 1725, Ap
14	US-10-060-756A-1726	Sequence 1726, Ap
14	US-10-060-756A-1727	Sequence 1727, Ap
19	US-10-205-309-284	Sequence 284, App
19	US-10-205-309-609	Sequence 609, App
20	US-10-087-325-15	Sequence 15, Appl
24	US-09-853-830-168	Sequence 168, App
24	US-09-934-489A-46	Sequence 46, Appl
24	US-10-438-729-167	Sequence 167, App
25	US-09-827-998-1102	Sequence 1102, Ap
25	US-09-827-998-1103	Sequence 1103, Ap
25	US-09-827-998-1104	Sequence 1104, Ap
25	US-09-827-998-1105	Sequence 1105, Ap
25	US-09-827-998-1106	Sequence 1106, Ap
25	US-09-827-998-1107	Sequence 1107, Ap
25	US-09-827-998-1108	Sequence 1108, Ap
25	US-09-827-998-1109	Sequence 1109, Ap
25	US-09-827-998-1110	Sequence 1110, Ap
25	US-09-827-998-1111	Sequence 1111, Ap
25	US-09-827-998-1112	Sequence 1112, Ap
25	US-09-827-998-1113	Sequence 1113, Ap
25	US-09-827-998-1114	Sequence 1114, Ap
25	US-09-827-998-1115	Sequence 1115, Ap
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25	US-10-215-112-7250	Sequence 7250, Ap
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25	US-10-098-263B-10333	Sequence 10333, A
25	US-10-098-263B-34049	Sequence 34049, A
25	US-10-098-263B-41737	Sequence 41737, A
25	US-10-098-263B-86720	Sequence 86720, A
25	US-10-098-263B-114427	Sequence 114427, A
25	US-10-098-263B-114428	Sequence 114428, A
25	US-10-098-263B-118057	Sequence 118057, A
25	US-10-098-263B-118058	Sequence 118058, A
25	US-10-098-263B-118243	Sequence 118243, A
25	US-10-098-263B-118244	Sequence 118244, A
25	US-10-098-263B-119613	Sequence 119613, A
33	US-09-765-272-359	Sequence 359, App
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38	US-09-780-533A-4693	Sequence 4693, App
38	US-09-877-478-4478	Sequence 4478, App
38	US-09-922-261-227	Sequence 227, App
42	US-10-032-585-1923	Sequence 1923, App
43	US-09-922-261-225	Sequence 225, App
53	US-09-997-931-40	Sequence 40, Appl
53	US-09-997-931-41	Sequence 41, Appl
59	US-10-085-906-282	Sequence 282, App
60	US-09-908-975-18382	Sequence 18382, A
60	US-09-908-975-31961	Sequence 31961, A
61	US-09-795-668-1332	Sequence 1332, Ap
61	US-09-795-668-1332	Sequence 1332, Ap
61	US-09-946-807-1332	Sequence 1332, Ap

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91	10	100.0	63	14	US-09-383-965-5096	Sequence 5096, App	164	14	100.0	178	10	US-10-092-154-327	Sequence 327, App
92	10	100.0	64	14	US-10-085-906-276	Sequence 276, App	165	10	100.0	181	11	US-09-754-853A-287	Sequence 287, App
93	10	100.0	65	12	US-09-308-975-28615	Sequence 27615, A	166	10	100.0	182	10	US-09-954-531-893	Sequence 893, App
94	10	100.0	65	12	US-09-308-975-28615	Sequence 28615, A	167	10	100.0	183	9	US-09-864-761-22682	Sequence 22682, A
95	10	100.0	65	12	US-10-032-585-135	Sequence 135, App	168	10	100.0	184	10	US-09-765-231A-48	Sequence 48, Appl
96	10	100.0	65	12	US-10-032-585-135	Sequence 135, App	169	10	100.0	186	9	US-09-864-761-31276	Sequence 31276, A
97	10	100.0	65	12	US-10-032-585-2790	Sequence 2790, App	170	10	100.0	186	12	US-09-803-668-5	Sequence 668, A
98	10	100.0	76	14	US-10-015-637-3	Sequence 3, Appl	171	10	100.0	186	12	US-10-243-475-13	Sequence 5, Appl
99	10	100.0	81	12	US-10-029-386-24504	Sequence 24504, A	172	10	100.0	189	10	US-09-923-261-213	Sequence 213, App
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101	10	100.0	87	9	US-09-864-761-29873	Sequence 29873, A	174	10	100.0	192	9	US-09-864-761-24940	Sequence 24940, A
102	10	100.0	88	12	US-10-029-386-24402	Sequence 24402, A	175	10	100.0	192	10	US-09-878-574-6130	Sequence 6130, App
103	10	100.0	89	9	US-09-864-761-31294	Sequence 31294, A	176	10	100.0	194	13	US-10-001-879-67	Sequence 67, Appl
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105	10	100.0	96	10	US-09-922-261-223	Sequence 223, App	178	10	100.0	198	10	US-09-765-231A-69	Sequence 69, Appl
106	10	100.0	97	9	US-09-864-761-25187	Sequence 25187, A	179	10	100.0	201	9	US-09-736-969A-80	Sequence 80, Appl
107	10	100.0	98	10	US-09-969-373-1521	Sequence 1521, App	180	10	100.0	201	10	US-09-878-574-9594	Sequence 9594, App
108	10	100.0	102	9	US-09-864-761-25485	Sequence 25485, A	181	10	100.0	201	10	US-09-974-300-4222	Sequence 4222, App
109	10	100.0	102	12	US-09-974-026-55	Sequence 55, Appl	182	10	100.0	201	12	US-10-238-075-1160	Sequence 1160, App
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117	10	100.0	118	12	US-10-029-386-15118	Sequence 15118, A	190	10	100.0	208	10	US-09-796-693-9479	Sequence 9479, App
118	10	100.0	118	14	US-10-007-280A-132	Sequence 132, App	191	10	100.0	208	14	US-10-040-863-9479	Sequence 9479, App
119	10	100.0	119	9	US-09-770-696-694	Sequence 694, App	192	10	100.0	210	9	US-09-864-761-27524	Sequence 27524, A
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122	10	100.0	121	11	US-09-818-875-198	Sequence 198, App	195	10	100.0	213	10	US-09-878-574-13095	Sequence 13095, A
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124	10	100.0	121	11	US-09-818-875-214	Sequence 214, App	197	10	100.0	215	12	US-10-287-274-45	Sequence 45, Appl
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126	10	100.0	121	11	US-09-818-875-318	Sequence 318, App	199	10	100.0	218	9	US-09-864-761-18972	Sequence 18972, A
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129	10	100.0	123	11	US-09-764-891-6108	Sequence 6108, App	202	10	100.0	225	14	US-10-040-863-9189	Sequence 9189, App
130	10	100.0	123	11	US-09-764-891-6109	Sequence 6109, App	203	10	100.0	226	9	US-09-563-817-279	Sequence 279, App
131	10	100.0	125	9	US-09-294-093B-56	Sequence 56, Appl	204	10	100.0	226	12	US-10-116-712-342	Sequence 342, App
132	10	100.0	125	9	US-09-563-817-780	Sequence 780, App	205	10	100.0	228	10	US-09-974-300-5635	Sequence 5635, App
133	10	100.0	126	10	US-09-922-261-219	Sequence 219, App	206	10	100.0	228	8	US-08-781-986A-4596	Sequence 4596, App
134	10	100.0	126	12	US-10-029-386-19178	Sequence 19178, A	207	10	100.0	230	14	US-10-103-524-560	Sequence 560, App
135	10	100.0	129	9	US-09-864-761-29992	Sequence 29992, A	208	10	100.0	231	9	US-09-864-761-21191	Sequence 21191, A
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137	10	100.0	132	11	US-09-962-967A-22	Sequence 22, Appl	210	10	100.0	231	10	US-09-796-692-3380	Sequence 3380, App
138	10	100.0	134	12	US-10-029-386-15576	Sequence 15576, A	211	10	100.0	231	14	US-10-040-863-1883	Sequence 1883, App
139	10	100.0	135	12	US-10-029-386-17260	Sequence 17260, A	212	10	100.0	232	14	US-10-060-036-1883	Sequence 1883, App
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145	10	100.0	150	10	US-09-969-373-201	Sequence 201, App	218	10	100.0	241	9	US-09-604-287A-360	Sequence 360, App
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150	10	100.0	156	10	US-09-974-300-4171	Sequence 4171, App	223	10	100.0	242	14	US-10-076-623-360	Sequence 360, App
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153	10	100.0	163	14	US-10-060-036-2004	Sequence 2004, App	226	10	100.0	250	9	US-09-294-093B-289	Sequence 289, App
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156	10	100.0	169	14	US-10-015-219-695	Sequence 695, App	229	10	100.0	254	12	US-09-814-353-16251	Sequence 16251, A
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158	10	100.0	174	9	US-09-864-761-29931	Sequence 29931, A	231	10	100.0	256	10	US-09-902-941-1077	Sequence 1077, App
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C 304	10	100.0	285	12	US-10-029-386-22548	Sequence 22548, A	C 377	10	100.0	319	12	US-09-942-052-726	Sequence 726, App
C 305	10	100.0	286	10	US-09-104-408-7	Sequence 7, Appl	C 378	10	100.0	320	9	US-09-759-143-224	Sequence 224, App
C 306	10	100.0	286	12	US-09-814-353-3363	Sequence 3363, App	C 379	10	100.0	320	9	US-09-780-669-224	Sequence 224, App
C 307	10	100.0	286	12	US-09-814-353-3684	Sequence 3684, App	C 380	10	100.0	320	9	US-09-030-606-224	Sequence 224, App
C 308	10	100.0	287	13	US-10-027-632-286859	Sequence 286859, A	C 381	10	100.0	320	9	US-09-822-827-224	Sequence 224, App

C 382	10	100.0	320	9	US-09-115-453-224	Sequence 224, App	455	10	100.0	346	12	US-09-873-319-233	Sequence 233, App
C 383	10	100.0	320	10	US-09-878-574-3072	Sequence 3072, Ap	456	10	100.0	346	12	US-09-960-706-386	Sequence 386, App
C 384	10	100.0	320	10	US-09-764-864-545	Sequence 545, App	457	10	100.0	346	14	US-10-228-794-81	Sequence 81, Appl
C 385	10	100.0	320	10	US-09-232-880-224	Sequence 224, App	458	10	100.0	346	14	US-10-106-698-2735	Sequence 2735, Ap
C 386	10	100.0	320	10	US-09-895-793-224	Sequence 224, App	459	10	100.0	347	10	US-09-867-701-10233	Sequence 10233, A
C 387	10	100.0	320	10	US-09-895-814-224	Sequence 224, App	460	10	100.0	347	12	US-10-116-719-614	Sequence 614, App
C 388	10	100.0	320	11	US-09-803-719-1955	Sequence 1955, Ap	c 461	10	100.0	347	14	US-10-060-036-1825	Sequence 1825, Ap
C 389	10	100.0	320	11	US-09-803-719-1955	Sequence 1955, Ap	c 461	10	100.0	347	14	US-10-060-036-1825	Sequence 1825, Ap
C 390	10	100.0	320	12	US-10-144-678A-224	Sequence 224, App	c 462	10	100.0	348	9	US-09-864-761-2092	Sequence 2092, Ap
C 391	10	100.0	320	12	US-09-814-353-4429	Sequence 4429, A	c 463	10	100.0	348	10	US-09-864-761-2092	Sequence 2092, Ap
C 392	10	100.0	320	12	US-10-294-025-224	Sequence 224, App	c 464	10	100.0	348	12	US-09-867-701-2327	Sequence 2327, Ap
C 393	10	100.0	320	12	US-10-012-896-224	Sequence 224, App	c 465	10	100.0	348	12	US-09-814-353-3169	Sequence 3169, Ap
C 394	10	100.0	320	14	US-10-010-940-224	Sequence 224, App	c 466	10	100.0	349	10	US-09-867-701-9513	Sequence 9513, Ap
C 395	10	100.0	321	11	US-09-910-009A-151	Sequence 151, App	c 467	10	100.0	349	10	US-09-974-300-3225	Sequence 3225, Ap
C 396	10	100.0	321	11	US-09-918-995-6771	Sequence 6771, Ap	c 468	10	100.0	349	11	US-09-764-891-1286	Sequence 1286, Ap
C 397	10	100.0	322	11	US-09-535-459-1981	Sequence 1981, Ap	c 469	10	100.0	349	13	US-10-040-733-889	Sequence 889, App
C 398	10	100.0	324	10	US-09-833-381-1209	Sequence 1209, Ap	470	10	100.0	350	9	US-09-770-791-705	Sequence 705, App
C 399	10	100.0	324	14	US-10-169-048-45	Sequence 45, Appl	471	10	100.0	350	13	US-10-027-632-54299	Sequence 54299, A
C 400	10	100.0	325	10	US-09-369-708-506	Sequence 506, App	472	10	100.0	350	13	US-10-027-632-180119	Sequence 180119, A
C 401	10	100.0	325	10	US-09-354-456-1667	Sequence 1667, App	c 473	10	100.0	351	10	US-09-867-701-854	Sequence 854, App
C 402	10	100.0	325	10	US-09-880-107-1838	Sequence 1838, Ap	474	10	100.0	353	10	US-09-560-863-669	Sequence 669, App
C 403	10	100.0	326	8	US-08-781-986A-4309	Sequence 4309, Ap	475	10	100.0	353	10	US-09-960-352-7434	Sequence 7434, Ap
C 404	10	100.0	326	11	US-09-910-009A-194	Sequence 194, App	476	10	100.0	353	10	US-09-983-965-155	Sequence 155, App
C 405	10	100.0	326	12	US-09-814-353-664	Sequence 664, App	c 477	10	100.0	353	11	US-09-953-999-35	Sequence 35, Appl
C 406	10	100.0	326	12	US-09-814-353-7040	Sequence 7040, Ap	478	10	100.0	353	11	US-09-803-719-945	Sequence 945, App
C 407	10	100.0	327	10	US-09-764-847-1130	Sequence 1130, Ap	479	10	100.0	353	11	US-09-803-719-1506	Sequence 1506, Ap
C 408	10	100.0	327	11	US-09-803-719-899	Sequence 899, App	c 480	10	100.0	354	9	US-09-815-242-9617	Sequence 9617, Ap
C 409	10	100.0	327	11	US-09-764-847-1130	Sequence 1130, Ap	c 481	10	100.0	354	10	US-09-764-847-80	Sequence 80, Appl
C 410	10	100.0	327	12	US-10-116-712-180	Sequence 180, App	c 482	10	100.0	354	10	US-09-938-842A-5319	Sequence 5319, Ap
C 411	10	100.0	327	14	US-10-092-154-1130	Sequence 1130, Ap	c 483	10	100.0	354	14	US-10-092-154-80	Sequence 80, Appl
C 412	10	100.0	328	10	US-09-764-847-1129	Sequence 1129, Ap	c 484	10	100.0	355	9	US-09-770-791-618	Sequence 618, App
C 413	10	100.0	328	14	US-10-092-154-1129	Sequence 1129, Ap	c 485	10	100.0	355	10	US-09-867-701-2400	Sequence 2400, Ap
C 414	10	100.0	329	11	US-09-803-719-878	Sequence 878, App	c 486	10	100.0	356	10	US-09-867-701-170	Sequence 170, App
C 415	10	100.0	330	10	US-09-867-701-278	Sequence 278, App	487	10	100.0	356	10	US-09-867-701-2002	Sequence 2002, Ap
C 416	10	100.0	330	14	US-10-083-357-254	Sequence 254, App	488	10	100.0	356	11	US-09-918-995-25782	Sequence 25782, A
C 417	10	100.0	331	13	US-10-027-632-28358	Sequence 28358, A	489	10	100.0	356	11	US-09-764-891-1552	Sequence 1552, Ap
C 418	10	100.0	332	12	US-10-029-386-24244	Sequence 24244, A	c 490	10	100.0	356	12	US-10-161-051-17	Sequence 17, Appl
C 419	10	100.0	332	12	US-10-116-712-446	Sequence 446, App	c 491	10	100.0	357	9	US-09-764-887-148	Sequence 148, App
C 420	10	100.0	332	12	US-10-116-712-446	Sequence 446, App	492	10	100.0	357	10	US-09-867-701-2408	Sequence 2408, Ap
C 421	10	100.0	333	10	US-09-867-701-110	Sequence 110, App	c 493	10	100.0	357	14	US-10-073-961-148	Sequence 148, App
C 422	10	100.0	333	12	US-09-814-353-4006	Sequence 4006, Ap	494	10	100.0	358	9	US-09-864-761-11220	Sequence 11220, A
C 423	10	100.0	333	12	US-09-814-353-10314	Sequence 10314, A	495	10	100.0	358	11	US-09-918-995-7467	Sequence 7467, Ap
C 424	10	100.0	333	12	US-09-814-353-10314	Sequence 10314, A	c 496	10	100.0	359	10	US-09-783-590-1751	Sequence 1751, Ap
C 425	10	100.0	334	9	US-09-814-353-19427	Sequence 19427, A	c 497	10	100.0	359	12	US-09-814-353-21980	Sequence 21980, A
C 426	10	100.0	334	11	US-09-783-590-1635	Sequence 1635, Ap	c 498	10	100.0	359	14	US-10-102-524-1534	Sequence 1534, Ap
C 427	10	100.0	334	11	US-09-918-995-18886	Sequence 18886, A	c 499	10	100.0	360	9	US-09-777-564-810	Sequence 810, App
C 428	10	100.0	335	9	US-09-784-423-26	Sequence 26, Appl	500	10	100.0	360	9	US-09-864-761-27334	Sequence 27334, A
C 429	10	100.0	335	10	US-09-983-965-3102	Sequence 3102, Ap							
C 430	10	100.0	335	10	US-09-796-692-6576	Sequence 6576, Ap							
C 431	10	100.0	335	13	US-10-027-632-288453	Sequence 288453, A							
C 432	10	100.0	335	14	US-10-040-862-6576	Sequence 6576, Ap							
C 433	10	100.0	336	10	US-09-974-300-8266	Sequence 8266, Ap							
C 434	10	100.0	336	12	US-10-238-075-555	Sequence 555, App							
C 435	10	100.0	337	10	US-09-929-493-7	Sequence 7, Appl							
C 436	10	100.0	337	11	US-09-803-719-1668	Sequence 1668, Ap							
C 437	10	100.0	337	11	US-09-918-995-19793	Sequence 19793, A							
C 438	10	100.0	338	12	US-10-270-487-7	Sequence 7, Appl							
C 439	10	100.0	338	11	US-09-803-719-963	Sequence 963, App							
C 440	10	100.0	339	13	US-10-027-632-128730	Sequence 128730, A							
C 441	10	100.0	339	14	US-10-060-036-1555	Sequence 1555, Ap							
C 442	10	100.0	339	14	US-10-313-542-168	Sequence 168, App							
C 443	10	100.0	340	10	US-09-920-300A-423	Sequence 423, App							
C 444	10	100.0	340	12	US-10-099-926-423	Sequence 423, App							
C 445	10	100.0	340	13	US-10-033-528-423	Sequence 423, App							
C 446	10	100.0	342	9	US-09-770-791-814	Sequence 814, App							
C 447	10	100.0	342	11	US-09-918-995-18543	Sequence 18543, A							
C 448	10	100.0	342	12	US-10-116-712-589	Sequence 589, App							
C 449	10	100.0	343	11	US-09-803-719-1667	Sequence 1667, Ap							
C 450	10	100.0	343	12	US-10-125-159-81	Sequence 81, Appl							
C 451	10	100.0	344	12	US-10-116-712-134	Sequence 134, App							
C 452	10	100.0	345	9	US-09-770-791-775	Sequence 775, App							
C 453	10	100.0	346	10	US-09-969-708-371	Sequence 371, App							
C 454	10	100.0	346	10	US-09-969-708-436	Sequence 436, App							

ALIGNMENTS

RESULT 1  
US-09-827-998-287/c  
; Sequence 287, Application US/09827998  
; Patent No. US20020102252A1  
; GENERAL INFORMATION:  
; APPLICANT: Gu, Yizhong  
; APPLICANT: Shannon, Mark  
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E  
; FILE REFERENCE: MDMORF-8  
; CURRENT APPLICATION NUMBER: US/09/827,998  
; CURRENT FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; NUMBER OF SEQ ID NOS: 1881  
; SOFTWARE: Aeonica Sequence Listing Engine  
; SEQ ID NO 287  
; LENGTH: 17  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-827-998-287



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Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 17 CTTCTCTTTT 8

RESULT 2
US-09-827-998-288/c
; Sequence 288, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWOF-8
; CURRENT APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 288
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-288

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 16 CTTCTCTTTT 7

RESULT 3
US-09-827-998-289/c
; Sequence 289, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWOF-8
; CURRENT APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 289
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-289

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 15 CTTCTCTTTT 6

RESULT 4
US-09-827-998-290/c
; Sequence 290, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWOF-8
; CURRENT APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 290
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-290

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 14 CTTCTCTTTT 5

RESULT 5
US-09-827-998-291/c
; Sequence 291, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWOF-8
; CURRENT APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 291
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-291

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 13 CTTCTCTTTT 4

RESULT 6
US-09-827-998-292/c
; Sequence 292, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
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; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 292
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-292

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3

RESULT 7
US-09-827-998-293/c
; Sequence 293, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 293
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-293

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 8
US-09-827-998-294/c
; Sequence 294, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 294
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-294

Query Match      100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3

RESULT 9
US-09-930-423-1740/c
; Sequence 1740, Application US/09930423
; Publication No. US20030092003A1
; GENERAL INFORMATION:
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: MEB00,918-A 400/027
; CURRENT APPLICATION NUMBER: US/09/930,423
; CURRENT FILING DATE: 2001-08-15
; NUMBER OF SEQ ID NOS: 4553
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1740
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-09-930-423-1740

Query Match      100.0%; Score 10; DB 11; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 10
US-09-930-423-1741/c
; Sequence 1741, Application US/09930423
; Publication No. US20030092003A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: MEB00,918-A 400/027
; CURRENT APPLICATION NUMBER: US/09/930,423
; CURRENT FILING DATE: 2001-08-15
; NUMBER OF SEQ ID NOS: 4553
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1741
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-09-930-423-1741

Query Match      100.0%; Score 10; DB 11; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3
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RESULT 11
US-09-745-237A-1740/c
; Sequence 1740, Application US/09745237A
; Publication No. US20030143708A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: 400/007 (MEHB00-918-A)
; CURRENT APPLICATION NUMBER: US/09/745,237A
; CURRENT FILING DATE: 2002-04-15
; NUMBER OF SEQ ID NOS: 4550
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1740
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo sapiens
US-09-745-237A-1740

Query Match          100.0%; Score 10; DB 12; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      14 CTTCTCTTTT 5

RESULT 12
US-09-745-237A-1741/c
; Sequence 1741, Application US/09745237A
; Publication No. US20030143708A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: 400/007 (MEHB00-918-A)
; CURRENT APPLICATION NUMBER: US/09/745,237A
; CURRENT FILING DATE: 2002-04-15
; NUMBER OF SEQ ID NOS: 4550
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1741
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo sapiens
US-09-745-237A-1741

Query Match          100.0%; Score 10; DB 12; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      12 CTTCTCTTTT 3

RESULT 13
US-10-060-756A-1720/c
; Sequence 1720, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1720
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1720/c

Query Match          100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      16 CTTCTCTTTT 7

RESULT 14
US-10-060-756A-1721/c
; Sequence 1721, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US/09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US/60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1721
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1721

Query Match          100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      17 CTTCTCTTTT 8

RESULT 15
US-10-060-756A-1722/c
```

```
; Sequence 1722, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1722
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1722

Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      15 CTTCTCTTTT 6

RESULT 16
US-10-060-756A-1723/c
; Sequence 1723, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1723
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
```

```
US-10-060-756A-1723
Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      14 CTTCTCTTTT 5

RESULT 17
US-10-060-756A-1724/c
; Sequence 1724, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1724
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1724

Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      13 CTTCTCTTTT 4

RESULT 18
US-10-060-756A-1725/c
; Sequence 1725, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
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; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aeomica Sequence Listing Engine
; SEQ ID NO 1725
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1725

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3

RESULT 19
US-10-060-756A-1726/c
; Sequence 1726, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aeomica Sequence Listing Engine
; SEQ ID NO 1726
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1726

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 20
US-10-060-756A-1727/c
; Sequence 1727, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
```

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; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aeomica Sequence Listing Engine
; SEQ ID NO 1727
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1727

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1

RESULT 21
US-10-205-309-284/c
; Sequence 284, Application US/10205309
; Publication No. US20030190635A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: RNA Interference Mediated Inhibition of Alzheimer's Disease Usin
; FILE REFERENCE: 900/033
; CURRENT APPLICATION NUMBER: US/10/205,309
; CURRENT FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 674
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 284
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Target sequence/siNA sense
US-10-205-309-284

Query Match 100.0%; Score 10; DB 12; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1

RESULT 22
US-10-205-309-609
; Sequence 609, Application US/10205309
; Publication No. US20030190635A1
; GENERAL INFORMATION:
```

```
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: RNA Interference Mediated Inhibition of Alzheimer's Disease Using
; TITLE OF INVENTION: Interfering RNA
; FILE REFERENCE: 900/033
; CURRENT APPLICATION NUMBER: US/10/205,309
; CURRENT FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 674
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 609
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: siNA antisense region
US-10-205-309-609

Query Match      100.0%; Score 10; DB 12; Length 19;
Best Local Similarity 30.0%; Pred. No. 1.6e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||:|:|:|:|
Db 10 CUUCUCUUU 19

RESULT 23
US-10-087-325-15/c
; Sequence 15, Application US/10087325
; Publication No. US20020192682A1
; GENERAL INFORMATION:
; APPLICANT: Escary, Jean-Louis
; TITLE OF INVENTION: NEW POLYNUCLEOTIDES AND POLYPEPTIDES OF THE IFNalpha-2 GENE
; FILE REFERENCE: 021349/0010
; CURRENT APPLICATION NUMBER: US/10/087,325
; CURRENT FILING DATE: 2002-03-01
; PRIOR APPLICATION NUMBER: FR 0102843
; PRIOR FILING DATE: 2001-03-01
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 15
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-087-325-15

Query Match      100.0%; Score 10; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||:|:|:|:|
Db 20 CTTCTCTTTT 11

RESULT 24
US-09-853-830-168/c
; Sequence 168, Application US/09853830
; Patent No. US20020107388A1
; GENERAL INFORMATION:
; APPLICANT: Vandenberg, Arthur A.
; TITLE OF INVENTION: Methods of Identifying and Monitoring
; TITLE OF INVENTION: Disease-Associated T Cells
; FILE REFERENCE: P-IM 4734
; CURRENT APPLICATION NUMBER: US/09/853,830
; CURRENT FILING DATE: 2001-09-18
; NUMBER OF SEQ ID NOS: 184
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 168
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-853-830-168
```

```
Query Match      100.0%; Score 10; DB 10; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||:|:|:|:|
Db 18 CTTCTCTTTT 9

RESULT 25
US-09-934-489A-46/c
; Sequence 46, Application US/09934489A
; Publication No. US20030108872A1
; GENERAL INFORMATION:
; APPLICANT: Sulavik, Mark
; APPLICANT: Ling, Losee Lucy
; APPLICANT: Opperman, Tlm
; APPLICANT: Moir, Don
; APPLICANT: Bunker, Christopher
; TITLE OF INVENTION: Genomics-Assisted Rapid Identification of Targets
; FILE REFERENCE: 032796-082
; CURRENT APPLICATION NUMBER: US/09/934,489A
; CURRENT FILING DATE: 2001-08-23
; PRIOR APPLICATION NUMBER: 2001-08-23,896
; PRIOR FILING DATE: 2000-08-23
; NUMBER OF SEQ ID NOS: 63
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: primer
US-09-934-489A-46

Query Match      100.0%; Score 10; DB 11; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||:|:|:|:|
Db 16 CTTCTCTTTT 7

RESULT 26
US-10-438-729-167/c
; Sequence 167, Application US/10438729
; Publication No. US20030190665A1
; GENERAL INFORMATION:
; APPLICANT: Vandenberg, Arthur
; TITLE OF INVENTION: METHODS OF SELECTING T CELL RECEPTOR V PEPTIDES FOR THERAPEUTIC
; FILE REFERENCE: 6915-65828
; CURRENT APPLICATION NUMBER: US/10/438,729
; CURRENT FILING DATE: 2003-05-14
; PRIOR APPLICATION NUMBER: 60/203,984
; PRIOR FILING DATE: 2000-05-12
; PRIOR APPLICATION NUMBER: 09/853,830
; PRIOR FILING DATE: 2001-05-10
; PRIOR APPLICATION NUMBER: 60/380,731
; PRIOR FILING DATE: 2002-05-14
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 167
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-438-729-167

Query Match      100.0%; Score 10; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
```

```
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 27
US-09-827-998-1102/c
; Sequence 1102, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1102
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1102

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 28
US-09-827-998-1103/c
; Sequence 1103, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1103
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1103

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 24 CTTCTCTTTT 15

RESULT 29
US-09-827-998-1104/c
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; Sequence 1104, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1104
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1104

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 23 CTTCTCTTTT 14

RESULT 30
US-09-827-998-1105/c
; Sequence 1105, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1105
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1105

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 22 CTTCTCTTTT 13

RESULT 31
US-09-827-998-1106/c
; Sequence 1106, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
```

```
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1106
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1106

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      21 CTTCTCTTTT 12

RESULT 32
US-09-827-998-1107/c
; Sequence 1107, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1107
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1107

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      20 CTTCTCTTTT 11

RESULT 33
US-09-827-998-1108/c
; Sequence 1108, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1108
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1108

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      18 CTTCTCTTTT 9

RESULT 34
US-09-827-998-1109/c
; Sequence 1109, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1109
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1109

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      18 CTTCTCTTTT 9

RESULT 35
US-09-827-998-1110/c
; Sequence 1110, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1110
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1110

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      19 CTTCTCTTTT 10
```



```
Db      17  CTTCTCTTTT 8

RESULT 36
US-09-827-998-1111/c
; Sequence 1111, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1111
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1111

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      16  CTTCTCTTTT 7

RESULT 37
US-09-827-998-1112/c
; Sequence 1112, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1112
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1112

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      15  CTTCTCTTTT 6

RESULT 38
US-09-827-998-1113/c
; Sequence 1113, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
```

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; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1113
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1113

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      14  CTTCTCTTTT 5

RESULT 39
US-09-827-998-1114/c
; Sequence 1114, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1114
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1114

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      13  CTTCTCTTTT 4

RESULT 40
US-09-827-998-1115/c
; Sequence 1115, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
```

; NUMBER OF SEQ ID NOS: 1881  
; SOFTWARE: Aescmca Sequence Listing Engine  
; SEQ ID NO 1115  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-827-998-1115

Query Match 100.0%; Score 10; DB 10; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 12 CTTCTCTTTT 3

RESULT 41  
US-09-827-998-1116/c  
; Sequence 1116, Application US/09827998  
; Patent No. US20020102252A1  
; GENERAL INFORMATION:  
; APPLICANT: Gu, Yizhong  
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E  
; FILE REFERENCE: MDMORF-8  
; CURRENT APPLICATION NUMBER: US/09/827,998  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; NUMBER OF SEQ ID NOS: 1881  
; SOFTWARE: Aescmca Sequence Listing Engine  
; SEQ ID NO 1116  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-827-998-1116

Query Match 100.0%; Score 10; DB 10; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 11 CTTCTCTTTT 2

RESULT 42  
US-09-827-998-1117/c  
; Sequence 1117, Application US/09827998  
; Patent No. US20020102252A1  
; GENERAL INFORMATION:  
; APPLICANT: Gu, Yizhong  
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E  
; FILE REFERENCE: MDMORF-8  
; CURRENT APPLICATION NUMBER: US/09/827,998  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; NUMBER OF SEQ ID NOS: 1881  
; SOFTWARE: Aescmca Sequence Listing Engine  
; SEQ ID NO 1117  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-827-998-1117

Query Match 100.0%; Score 10; DB 10; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 10 CTTCTCTTTT 1

RESULT 43  
US-10-060-756A-3717/c  
; Sequence 3717, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; CURRENT FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Aescmca Sequence Listing Engine  
; SEQ ID NO 3717  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3717

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 25 CTTCTCTTTT 16

RESULT 44  
US-10-060-756A-3718/c  
; Sequence 3718, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; CURRENT FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30

```
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3718
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3718
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      24 CTTCTCTTTT 15
```

```
RESULT 45
US-10-060-756A-3719/c
; Sequence 3719, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3719
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3719
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      23 CTTCTCTTTT 14
```

```
RESULT 46
US-10-060-756A-3720/c
; Sequence 3720, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
```

```
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3720
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3720
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      22 CTTCTCTTTT 13
```

```
RESULT 47
US-10-060-756A-3721/c
; Sequence 3721, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3721
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3721
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

Db 21 CTTCTCTTTT 12  
|||||

## RESULT 48

US-10-060-756A-3722/c  
; Sequence 3722, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; PRIOR FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Acomica Sequence Listing Engine  
; SEQ ID NO 3722  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3722

Query Match 100.0%; Score 10; DB 14; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||

Db 20 CTTCTCTTTT 11  
|||||

## RESULT 49

US-10-060-756A-3723/c  
; Sequence 3723, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; PRIOR FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09

; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Acomica Sequence Listing Engine  
; SEQ ID NO 3723  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3723

Query Match 100.0%; Score 10; DB 14; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||

Db 19 CTTCTCTTTT 10  
|||||

## RESULT 50

US-10-060-756A-3724/c  
; Sequence 3724, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; PRIOR FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Acomica Sequence Listing Engine  
; SEQ ID NO 3724  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3724

Query Match 100.0%; Score 10; DB 14; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|||||

Db 18 CTTCTCTTTT 9  
|||||

## RESULT 51

US-10-060-756A-3725/c  
; Sequence 3725, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; PRIOR FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664

```
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3725
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3725
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
      |||||
Db 17 CTTCTCTTTT 8
```

```
RESULT 52
US-10-060-756A-3726/c
/ Sequence 3726, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3726
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3726
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
      |||||
Db 16 CTTCTCTTTT 7
```

```
RESULT 53
US-10-060-756A-3727/c
/ Sequence 3727, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3727
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3727
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
      |||||
Db 15 CTTCTCTTTT 6
```

```
RESULT 54
US-10-060-756A-3728/c
/ Sequence 3728, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3728
/ LENGTH: 25
```

; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3728

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
||| |||||  
Db 14 CTTCTCTTTT 5

## RESULT 55

US-10-060-756A-3729/c  
; Sequence 3729, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A  
; CURRENT FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Aecomica Sequence Listing Engine  
; SEQ ID NO 3729  
; LENGTH: 25  
; TYPE: DNA

; ORGANISM: Homo sapiens  
US-10-060-756A-3729

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
||| |||||  
Db 13 CTTCTCTTTT 4

## RESULT 56

US-10-060-756A-3730/c  
; Sequence 3730, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian  
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A  
; CURRENT FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665

; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Aecomica Sequence Listing Engine  
; SEQ ID NO 3730  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3730

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
||| |||||  
Db 12 CTTCTCTTTT 3

## RESULT 57

US-10-060-756A-3731/c  
; Sequence 3731, Application US/10060756A  
; Publication No. US20030046717A1  
; GENERAL INFORMATION:  
; APPLICANT: Zhang, Jian

; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN  
; FILE REFERENCE: PB0177  
; CURRENT APPLICATION NUMBER: US/10/060,756A  
; CURRENT FILING DATE: 2002-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/327,898  
; PRIOR FILING DATE: 2001-10-09  
; NUMBER OF SEQ ID NOS: 4804  
; SOFTWARE: Aecomica Sequence Listing Engine  
; SEQ ID NO 3731  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-060-756A-3731

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
||| |||||  
Db 11 CTTCTCTTTT 2

## RESULT 58

US-10-060-756A-3732/c  
; Sequence 3732, Application US/10060756A  
; Publication No. US20030046717A1

```
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aeomica Sequence Listing Engine
; SEQ ID NO 3732
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3732

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      10 CTTCTCTTTT 1

RESULT 59
US-10-215-112-7249/c
; Sequence 7249, Application US/10215112
; Publication No. US20030082596A1
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: 3119
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7249
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7249

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      13 CTTCTCTTTT 4

RESULT 60
US-10-215-112-7250/c
; Sequence 7250, Application US/10215112
; Publication No. US20030082596A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: 3119
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7250
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7250

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      12 CTTCTCTTTT 3

RESULT 61
US-10-215-112-7372/c
; Sequence 7372, Application US/10215112
; Publication No. US20030082596A1
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: 3119
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7372
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7372

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      12 CTTCTCTTTT 3

RESULT 62
US-10-098-263B-6553
; Sequence 6553, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6553
; LENGTH: 25
; TYPE: DNA
```

```

; ORGANISM: Homo sapien
US-10-098-263B-6553

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 7 CTTCTCTTTT 16

RESULT 63
US-10-098-263B-10333
; Sequence 10333, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 10333
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-10333

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 64
US-10-098-263B-86720/c
; Sequence 86720, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 86720
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-86720

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 5 CTTCTCTTTT 14

RESULT 65
US-10-098-263B-34049
; Sequence 34049, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 34049
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-34049

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 20

RESULT 66
US-10-098-263B-114427
; Sequence 114427, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 114427
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-41737
; Sequence 41737, Application US/10098263B

```



## US-10-098-263B-114427

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 16 CTTCTCTTTT 25

## RESULT 68

US-10-098-263B-114428  
; Sequence 114428, Application US/10098263B  
; Publication No. US20030104410A1  
; GENERAL INFORMATION:  
; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 114428  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-114428

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 16 CTTCTCTTTT 25

## RESULT 69

US-10-098-263B-118057  
; Sequence 118057, Application US/10098263B  
; Publication No. US20030104410A1  
; GENERAL INFORMATION:  
; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 118057  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-118057

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 15 CTTCTCTTTT 24

## RESULT 70

US-10-098-263B-118058  
; Sequence 118058, Application US/10098263B  
; Publication No. US20030104410A1

## ; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 118058  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-118058

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 15 CTTCTCTTTT 24

## RESULT 71

US-10-098-263B-118243  
; Sequence 118243, Application US/10098263B  
; Publication No. US20030104410A1  
; GENERAL INFORMATION:  
; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 118243  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-118243

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 15 CTTCTCTTTT 24

## RESULT 72

US-10-098-263B-118244  
; Sequence 118244, Application US/10098263B  
; Publication No. US20030104410A1  
; GENERAL INFORMATION:  
; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 118244  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-118244

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 15 CTTCTCTTTT 24

## RESULT 73

US-10-098-263B-119613  
; Sequence 119613, Application US/10098263B  
; Publication No. US20030104410A1  
; GENERAL INFORMATION:  
; APPLICANT: Mittman, Michael  
; TITLE OF INVENTION: Human Microarray  
; FILE REFERENCE: 3118.1  
; CURRENT APPLICATION NUMBER: US/10/098,263B  
; CURRENT FILING DATE: 2003-01-08  
; PRIOR APPLICATION NUMBER: 60/276,759  
; PRIOR FILING DATE: 2001-03-16  
; NUMBER OF SEQ ID NOS: 131066  
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1  
; SEQ ID NO 119613  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-10-098-263B-119613

Query Match 100.0%; Score 10; DB 14; Length 25;  
Best Local Similarity 100.0%; Pred. No. 1.6e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 13 CTTCTCTTTT 22

## RESULT 74

US-09-765-272-359/c  
; Sequence 359, Application US/09765272  
; Patent No. US20020061545A1  
; GENERAL INFORMATION:  
; APPLICANT: Choi et. al.  
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines  
; NUMBER OF SEQUENCES: 452  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Human Genome Sciences, Inc.  
; STREET: 9410 Key West Avenue  
; CITY: Rockville  
; STATE: Maryland  
; COUNTRY: USA  
; ZIP: 20850  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage  
; COMPUTER: HP Vectra 486/33  
; OPERATING SYSTEM: MSDOS version 6.2  
; SOFTWARE: ASCII Text  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/765,272  
; FILING DATE: 22-Jan-2001  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/961,083  
; FILING DATE: <Unknown>  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Brookes, A. Anders  
; REGISTRATION NUMBER: 36,373  
; REFERENCE/DOCKET NUMBER: PB340P2  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (301) 309-8504  
; TELEFAX: (301) 309-8512

; INFORMATION FOR SEQ ID NO: 359:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 33 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:  
US-09-765-272-359

Query Match 100.0%; Score 10; DB 9; Length 33;  
Best Local Similarity 100.0%; Pred. No. 1.5e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 23 CTTCTCTTTT 14

## RESULT 75

US-09-780-533A-2783/c  
; Sequence 2783, Application US/09780533A  
; Publication No. US20030060611A1  
; GENERAL INFORMATION:  
; APPLICANT: Ribozyme Pharmaceuticals, Inc.  
; APPLICANT: Blatt, Larry  
; APPLICANT: McSwiggen, Jim  
; APPLICANT: Chowrira, Bharat  
; APPLICANT: Haerberli, Pete  
; TITLE OF INVENTION: Method and Reagent for the Inhibition of NOGO Gene  
; FILE REFERENCE: MBH00,878-A (400/011)  
; CURRENT APPLICATION NUMBER: US/09/780,533A  
; CURRENT FILING DATE: 2001-02-09  
; PRIOR APPLICATION NUMBER: US 60/181,797  
; PRIOR FILING DATE: 2000-02-11  
; NUMBER OF SEQ ID NOS: 6679  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 2783  
; LENGTH: 38  
; TYPE: RNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid  
US-09-780-533A-2783

Query Match 100.0%; Score 10; DB 11; Length 38;  
Best Local Similarity 100.0%; Pred. No. 1.5e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
| | | | |  
Db 38 CTTCTCTTTT 29

## RESULT 76

US-09-780-533A-4693  
; Sequence 4693, Application US/09780533A  
; Publication No. US20030060611A1  
; GENERAL INFORMATION:  
; APPLICANT: Ribozyme Pharmaceuticals, Inc.  
; APPLICANT: Blatt, Larry  
; APPLICANT: McSwiggen, Jim  
; APPLICANT: Chowrira, Bharat  
; APPLICANT: Haerberli, Pete  
; TITLE OF INVENTION: Method and Reagent for the Inhibition of NOGO Gene  
; FILE REFERENCE: MBH00,878-A (400/011)  
; CURRENT APPLICATION NUMBER: US/09/780,533A  
; CURRENT FILING DATE: 2001-02-09  
; PRIOR APPLICATION NUMBER: US 60/181,797  
; PRIOR FILING DATE: 2000-02-11  
; NUMBER OF SEQ ID NOS: 6679  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 4693  
; LENGTH: 38

; TYPE: RNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid  
US-09-780-533A-4693

Query Match 100.0%; Score 10; DB 11; Length 38;  
Best Local Similarity 30.0%; Pred. No. 1.5e+04;  
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|::|  
Db 29 CUUCUCUUU 38

## RESULT 77

US-09-877-478-4478  
; Sequence 4478, Application US/09877478  
; Publication No. US20030068301A1  
; GENERAL INFORMATION:  
; APPLICANT: Ribozyme Pharmaceuticals, Inc.  
; APPLICANT: Draper, Kenneth  
; APPLICANT: Blatt, Larry  
; APPLICANT: McSwiggen, Jim  
; APPLICANT: Morrissey, Dave  
; TITLE OF INVENTION: Method and Reagent for Inhibiting Hepatitis B Virus Replication  
; FILE REFERENCE: MBH00-845-H (400/029)  
; CURRENT APPLICATION NUMBER: US/09/877,478  
; CURRENT FILING DATE: 2001-12-31  
; PRIOR APPLICATION NUMBER: US 07/882,712  
; PRIOR FILING DATE: 1992-05-14  
; PRIOR APPLICATION NUMBER: US 09/531,025  
; PRIOR FILING DATE: 2000-03-20  
; PRIOR APPLICATION NUMBER: US 09/636,385  
; PRIOR FILING DATE: 2000-08-09  
; PRIOR APPLICATION NUMBER: US 09/696,347  
; PRIOR FILING DATE: 2000-10-24  
; PRIOR APPLICATION NUMBER: US 08/193,627  
; PRIOR FILING DATE: 1994-02-07  
; PRIOR APPLICATION NUMBER: US 08/433,993  
; PRIOR FILING DATE: 1995-05-04  
; PRIOR APPLICATION NUMBER: US 08/434,504  
; PRIOR FILING DATE: 1995-05-04  
; PRIOR APPLICATION NUMBER: US 09/436,430  
; PRIOR FILING DATE: 1999-11-08  
; NUMBER OF SEQ ID NOS: 6586  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 4478  
; LENGTH: 38  
; TYPE: RNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid  
NAME/KEY: misc feature  
LOCATION: (11)..(16)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
NAME/KEY: misc feature  
LOCATION: (20)..(20)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
NAME/KEY: misc feature  
LOCATION: (22)..(22)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
NAME/KEY: misc feature  
LOCATION: (24)..(25)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
NAME/KEY: misc feature  
LOCATION: (28)..(28)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
NAME/KEY: misc feature  
LOCATION: (30)..(30)  
OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage  
US-09-877-478-4478

Query Match 100.0%; Score 10; DB 11; Length 38;  
Best Local Similarity 30.0%; Pred. No. 1.5e+04;  
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|::|  
Db 29 CUUCUCUUU 38

## RESULT 78

US-09-792-818-944/c  
; Sequence 944, Application US/09792818  
; Publication No. US20030134806A1  
; GENERAL INFORMATION:  
; APPLICANT: Ribozyme Pharmaceuticals, Inc.  
; APPLICANT: Jarvis, Thale  
; APPLICANT: Von Carlowitz, Ira  
; APPLICANT: McSwiggen, Jim  
; APPLICANT: Hamblin, Paul  
; APPLICANT: Ellis, Jonathan  
; TITLE OF INVENTION: Method and Reagent for the Inhibition of Grb-2-related with Inse  
; FILE REFERENCE: MBH00-901-A (400/013)  
; CURRENT APPLICATION NUMBER: US/09/792,818  
; CURRENT FILING DATE: 2001-02-23  
; NUMBER OF SEQ ID NOS: 2304  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 944  
; LENGTH: 38  
; TYPE: RNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid  
US-09-792-818-944

Query Match 100.0%; Score 10; DB 12; Length 38;  
Best Local Similarity 100.0%; Pred. No. 1.5e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|::|  
Db 38 CTTCTCTTTT 29

## RESULT 79

US-09-922-261-227/c  
; Sequence 227, Application US/09922261  
; Patent No. US20020111471A1  
; GENERAL INFORMATION:  
; APPLICANT: COGENT NEUROSCIENCE, Inc.  
; APPLICANT: Lo, Donald C.  
; APPLICANT: Barney, Shawn  
; APPLICANT: Thomas, Mary Beth  
; APPLICANT: Portbury, Stuart D.  
; APPLICANT: Putnam, Kasturi  
; APPLICANT: Katz, Lawrence C.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING  
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING  
; FILE REFERENCE: 10001-005-999  
; CURRENT APPLICATION NUMBER: US/09/922,261  
; CURRENT FILING DATE: 2001-08-03  
; PRIOR APPLICATION NUMBER: US/09/461,697  
; PRIOR FILING DATE: 1999-12-14  
; NUMBER OF SEQ ID NOS: 466  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 227  
; LENGTH: 42  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-922-261-227

Query Match 100.0%; Score 10; DB 10; Length 42;

```
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 80
US-10-032-585-1923
; Sequence 1923, Application US/10032585
; Publication No. US20030180953A1
; GENERAL INFORMATION:
; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1923
; LENGTH: 43
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-1923

Query Match 100.0%; Score 10; DB 12; Length 43;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 21

RESULT 81
US-09-922-261-225/c
; Sequence 225, Application US/09922261
; Patent No. US20020111471A1
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/922,261
; CURRENT FILING DATE: 2001-08-03
; PRIOR APPLICATION NUMBER: US/09/461,697
; PRIOR FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 225
; LENGTH: 48
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-922-261-225

Query Match 100.0%; Score 10; DB 10; Length 48;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 20 CTTCTCTTTT 11

Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
RESULT 82
US-09-997-931-40/c
; Sequence 40, Application US/09997931
; Publication No. US20030087241A1
; GENERAL INFORMATION:
; APPLICANT: University of Rochester
; APPLICANT: Kool, Eric
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND DNA
; FILE REFERENCE: 220.00010142
; CURRENT APPLICATION NUMBER: US/09/997,931
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: US 09/569,344
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 08/805,631
; PRIOR FILING DATE: 1997-02-26
; PRIOR APPLICATION NUMBER: US 08/393,439
; PRIOR FILING DATE: 1995-02-23
; PRIOR APPLICATION NUMBER: US 08/047,860
; NUMBER OF SEQ ID NOS: 129
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 40
; LENGTH: 53
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: 53mer circle
US-09-997-931-40

Query Match 100.0%; Score 10; DB 11; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 83
US-09-997-931-41
; Sequence 41, Application US/09997931
; Publication No. US20030087241A1
; GENERAL INFORMATION:
; APPLICANT: University of Rochester
; APPLICANT: Kool, Eric
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND DNA
; FILE REFERENCE: 220.00010142
; CURRENT APPLICATION NUMBER: US/09/997,931
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: US 09/569,344
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 08/805,631
; PRIOR FILING DATE: 1997-02-26
; PRIOR APPLICATION NUMBER: US 08/393,439
; PRIOR FILING DATE: 1995-02-23
; PRIOR APPLICATION NUMBER: US 08/047,860
; NUMBER OF SEQ ID NOS: 129
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 41
; LENGTH: 53
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: stem-loop multimer which binds HIV-1 gag RNA
US-09-997-931-41

Query Match 100.0%; Score 10; DB 11; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.5e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;
```

QY 1 CTTCTCTTTT 10  
|::|::|::|  
Db 38 CUUCUCUUU 47

RESULT 84  
US-10-085-906-282  
; Sequence 282, Application US/10085906  
; Publication No. US20030054371A1  
; GENERAL INFORMATION:  
; APPLICANT: Ying, Vincent  
; APPLICANT: Wu, Paul  
; APPLICANT: Gray, Gary S.  
; TITLE OF INVENTION: POLYMORPHIC ELEMENTS IN THE  
; TITLE OF INVENTION: COSTIMULATORY RECEPTOR LOCUS AND USES THEREOF  
; FILE REFERENCE: GNN-5343CP2  
; CURRENT APPLICATION NUMBER: US/10/085,906  
; CURRENT FILING DATE: 2002-02-27  
; PRIOR APPLICATION NUMBER: US 60/126,215  
; PRIOR FILING DATE: 1999-03-25  
; PRIOR APPLICATION NUMBER: US 09/534,061  
; PRIOR FILING DATE: 2000-03-24  
; PRIOR APPLICATION NUMBER: PCT/US00/07938  
; PRIOR FILING DATE: 2000-03-24  
; NUMBER OF SEQ ID NOS: 545  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 282  
; LENGTH: 59  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-085-906-282

Query Match 100.0%; Score 10; DB 14; Length 59;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|  
Db 45 CTTCTCTTTT 54

RESULT 85  
US-09-908-975-18382  
; Sequence 18382, Application US/09908975  
; Publication No. US20030165843A1  
; GENERAL INFORMATION:  
; APPLICANT: SHOSHAN, Avi  
; APPLICANT: WASSERMAN, Alon  
; APPLICANT: MINTZ, Eli  
; APPLICANT: MINTZ, Liat  
; APPLICANT: FAIGLER, Simchon  
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICE  
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME  
; FILE REFERENCE: 36688-0005  
; CURRENT APPLICATION NUMBER: US/09/908,975  
; CURRENT FILING DATE: 2001-07-20  
; PRIOR APPLICATION NUMBER: US 60/287,724  
; PRIOR FILING DATE: 2001-05-02  
; PRIOR APPLICATION NUMBER: US 60/221,607  
; PRIOR FILING DATE: 2000-07-28  
; NUMBER OF SEQ ID NOS: 32337  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 18382  
; LENGTH: 60  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-908-975-18382

Query Match 100.0%; Score 10; DB 12; Length 60;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 43 CTTCTCTTTT 52  
|::|::|::|

RESULT 86  
US-09-908-975-31961/c  
; Sequence 31961, Application US/09908975  
; Publication No. US20030165843A1  
; GENERAL INFORMATION:  
; APPLICANT: SHOSHAN, Avi  
; APPLICANT: WASSERMAN, Alon  
; APPLICANT: MINTZ, Eli  
; APPLICANT: MINTZ, Liat  
; APPLICANT: FAIGLER, Simchon  
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICE  
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME  
; FILE REFERENCE: 36688-0005  
; CURRENT APPLICATION NUMBER: US/09/908,975  
; CURRENT FILING DATE: 2001-07-20  
; PRIOR APPLICATION NUMBER: US 60/287,724  
; PRIOR FILING DATE: 2001-05-02  
; PRIOR APPLICATION NUMBER: US 60/221,607  
; PRIOR FILING DATE: 2000-07-28  
; NUMBER OF SEQ ID NOS: 32337  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 31961  
; LENGTH: 60  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-908-975-31961

Query Match 100.0%; Score 10; DB 12; Length 60;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|  
Db 48 CTTCTCTTTT 39

RESULT 87  
US-09-795-668-1332  
; Sequence 1332, Application US/09795668  
; Patent No. US20020045577A1  
; GENERAL INFORMATION:  
; APPLICANT: Stefansson, Hreinn  
; APPLICANT: Steinhorsdottir, Valgerdur  
; APPLICANT: Gulcher, Jeffrey R.  
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE  
; FILE REFERENCE: 2345-2004-001  
; CURRENT APPLICATION NUMBER: US/09/795,668  
; CURRENT FILING DATE: 2001-02-28  
; PRIOR APPLICATION NUMBER: US 09/515,716  
; PRIOR FILING DATE: 2000-02-28  
; NUMBER OF SEQ ID NOS: 1531  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1332  
; LENGTH: 61  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-795-668-1332

Query Match 100.0%; Score 10; DB 9; Length 61;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10  
|::|::|::|  
Db 17 CTTCTCTTTT 26

RESULT 88  
US-09-795-686-1332

; Sequence 1332, Application US/09795686  
; Patent No. US2002009495A1  
; GENERAL INFORMATION:  
; APPLICANT: Stefansson, Hreinn  
; APPLICANT: Steinhorsdottir, Valgerdur  
; APPLICANT: Gulcher, Jeffrey R.  
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE  
; FILE REFERENCE: 2345.2005-001  
; CURRENT APPLICATION NUMBER: US/09/795,686  
; CURRENT FILING DATE: 2001-02-28  
; PRIOR APPLICATION NUMBER: US 09/515,715  
; PRIOR FILING DATE: 2000-02-28  
; NUMBER OF SEQ ID NOS: 1531  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1332  
; LENGTH: 61  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-795-686-1332

Query Match 100.0%; Score 10; DB 9; Length 61;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 17 CTTCTCTTTT 26

## RESULT 89

US-09-946-807-1332  
; Sequence 1332, Application US/09946807  
; Patent No. US20020165144A1  
; GENERAL INFORMATION:  
; APPLICANT: Stefansson, Hreinn  
; APPLICANT: Steinhorsdottir, Valgerdur  
; APPLICANT: Gulcher, Jeffrey R.  
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE  
; FILE REFERENCE: 2345.2004-001  
; CURRENT APPLICATION NUMBER: US/09/946,807  
; CURRENT FILING DATE: 2001-09-05  
; PRIOR APPLICATION NUMBER: US/09/795,668  
; PRIOR FILING DATE: 2001-02-28  
; PRIOR APPLICATION NUMBER: US 09/515,716  
; PRIOR FILING DATE: 2000-02-28  
; NUMBER OF SEQ ID NOS: 1531  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1332  
; LENGTH: 61  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-946-807-1332

Query Match 100.0%; Score 10; DB 10; Length 61;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 17 CTTCTCTTTT 26

## RESULT 90

US-10-027-632-58348  
; Sequence 58348, Application US/10027632  
; GENERAL INFORMATION:  
; APPLICANT: Wang, David G.  
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
; POLYMORPHISMS IN THE HUMAN GENOME  
; FILE REFERENCE: 108827.129  
; CURRENT APPLICATION NUMBER: US/10/027,632  
; CURRENT FILING DATE: 2002-04-30  
; PRIOR APPLICATION NUMBER: US 60/218,006

; PRIOR FILING DATE: 2000-07-12  
; PRIOR APPLICATION NUMBER: US 60/198,676  
; PRIOR FILING DATE: 2000-04-20  
; PRIOR APPLICATION NUMBER: US 60/193,483  
; PRIOR FILING DATE: 2000-03-29  
; PRIOR APPLICATION NUMBER: US 60/185,218  
; PRIOR FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: US 60/167,363  
; PRIOR FILING DATE: 1999-11-23  
; PRIOR APPLICATION NUMBER: US 60/156,358  
; PRIOR FILING DATE: 1999-09-28  
; PRIOR APPLICATION NUMBER: US 60/146,002  
; PRIOR FILING DATE: 1999-08-09  
; NUMBER OF SEQ ID NOS: 325720  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 58348  
; LENGTH: 61  
; TYPE: DNA  
; ORGANISM: Human  
US-10-027-632-58348

Query Match 100.0%; Score 10; DB 13; Length 61;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 16 CTTCTCTTTT 25

## RESULT 91

US-09-983-965-5096  
; Sequence 5096, Application US/09983965  
; Patent No. US20020137160A1  
; GENERAL INFORMATION:  
; APPLICANT: Warren, Wesley C.  
; APPLICANT: Tao, Nengbing  
; APPLICANT: Byatt, John C.  
; APPLICANT: Mathialagan, Nagappan  
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND  
; FILE REFERENCE: 37-21(10297)C  
; CURRENT APPLICATION NUMBER: US/09/983,965  
; CURRENT FILING DATE: 2001-10-26  
; PRIOR APPLICATION NUMBER: US 09/465,231  
; PRIOR FILING DATE: 1999-12-15  
; PRIOR APPLICATION NUMBER: US 60/113,678  
; PRIOR FILING DATE: 1998-12-17  
; NUMBER OF SEQ ID NOS: 5912  
; SEQ ID NO 5096  
; LENGTH: 63  
; TYPE: DNA  
; ORGANISM: Bos taurus  
; FEATURE:  
; OTHER INFORMATION: Clone ID: 31-LIB34-084-Q1-E1-H11  
US-09-983-965-5096

Query Match 100.0%; Score 10; DB 10; Length 63;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||  
Db 24 CTTCTCTTTT 33

## RESULT 92

US-10-085-906-276/c  
; Sequence 276, Application US/10085906  
; Publication No. US20030054371A1  
; GENERAL INFORMATION:  
; APPLICANT: Ying, Vincent  
; APPLICANT: Wu, Paul

; APPLICANT: Gray, Gary S.  
; TITLE OF INVENTION: POLYMORPHIC ELEMENTS IN THE  
; FILE REFERENCE: GNN-5343CP2  
; CURRENT APPLICATION NUMBER: US/10/085,906  
; CURRENT FILING DATE: 2002-02-27  
; PRIOR APPLICATION NUMBER: US 60/126,215  
; PRIOR FILING DATE: 1999-03-25  
; PRIOR APPLICATION NUMBER: US 09/534,061  
; PRIOR FILING DATE: 2000-03-24  
; PRIOR APPLICATION NUMBER: PCT/US00/07938  
; PRIOR FILING DATE: 2000-03-24  
; NUMBER OF SEQ ID NOS: 545  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 276  
; LENGTH: 64  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-085-906-276

Query Match 100.0%; Score 10; DB 14; Length 64;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 43 CTTCTCTTTT 34

RESULT 93  
US-09-908-975-27615  
; Sequence 27615, Application US/09908975  
; Publication No. US20030165843A1  
; GENERAL INFORMATION:  
; APPLICANT: SHOSHAN, Avi  
; APPLICANT: WASSERMAN, Alon  
; APPLICANT: MINTZ, Eli  
; APPLICANT: MINTZ, Liat  
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING  
; FILE REFERENCE: 36688-0005  
; CURRENT APPLICATION NUMBER: US/09/908,975  
; CURRENT FILING DATE: 2001-07-20  
; PRIOR APPLICATION NUMBER: US 60/287,724  
; PRIOR FILING DATE: 2001-05-02  
; PRIOR APPLICATION NUMBER: US 60/221,607  
; PRIOR FILING DATE: 2000-07-28  
; NUMBER OF SEQ ID NOS: 32337  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 27615  
; LENGTH: 65  
; TYPE: DNA  
; ORGANISM: Mus musculus  
US-09-908-975-27615

Query Match 100.0%; Score 10; DB 12; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 47 CTTCTCTTTT 56

RESULT 94  
US-09-908-975-28061/c  
; Sequence 28061, Application US/09908975  
; Publication No. US20030165843A1  
; GENERAL INFORMATION:  
; APPLICANT: SHOSHAN, Avi  
; APPLICANT: WASSERMAN, Alon  
; APPLICANT: MINTZ, Eli

; APPLICANT: MINTZ, Liat  
; APPLICANT: FAIGLER, Simchon  
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING  
; FILE REFERENCE: 36688-0005  
; CURRENT APPLICATION NUMBER: US/09/908,975  
; CURRENT FILING DATE: 2001-07-20  
; PRIOR APPLICATION NUMBER: US 60/287,724  
; PRIOR FILING DATE: 2001-05-02  
; PRIOR APPLICATION NUMBER: US 60/221,607  
; PRIOR FILING DATE: 2000-07-28  
; NUMBER OF SEQ ID NOS: 32337  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 28061  
; LENGTH: 65  
; TYPE: DNA  
; ORGANISM: Mus musculus  
US-09-908-975-28061

Query Match 100.0%; Score 10; DB 12; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 41 CTTCTCTTTT 32

RESULT 95  
US-10-032-585-135  
; Sequence 135, Application US/10032585  
; Publication No. US20030180953A1  
; GENERAL INFORMATION:  
; APPLICANT: Terry, Roemer D.  
; APPLICANT: Bo, Jiang  
; APPLICANT: Charles, Boone  
; APPLICANT: Howard, Bussey  
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery  
; FILE REFERENCE: 10182-005-999  
; CURRENT APPLICATION NUMBER: US/10/032,585  
; CURRENT FILING DATE: 2001-12-20  
; NUMBER OF SEQ ID NOS: 8000  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 135  
; LENGTH: 65  
; TYPE: DNA  
; ORGANISM: Candida albicans  
US-10-032-585-135

Query Match 100.0%; Score 10; DB 12; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CTTCTCTTTT 10  
Db 2 CTTCTCTTTT 11

RESULT 96  
US-10-032-585-152  
; Sequence 152, Application US/10032585  
; Publication No. US20030180953A1  
; GENERAL INFORMATION:  
; APPLICANT: Terry, Roemer D.  
; APPLICANT: Bo, Jiang  
; APPLICANT: Charles, Boone  
; APPLICANT: Howard, Bussey  
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery  
; FILE REFERENCE: 10182-005-999  
; CURRENT APPLICATION NUMBER: US/10/032,585  
; CURRENT FILING DATE: 2001-12-20  
; NUMBER OF SEQ ID NOS: 8000  
; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 152  
; LENGTH: 65  
; TYPE: DNA  
; ORGANISM: Candida albicans  
US-10-032-585-152

Query Match 100.0%; Score 10; DB 12; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 15 CTTCTCTTTT 24

## RESULT 97

US-10-032-585-2790  
; Sequence 2790, Application US/10032585  
; Publication No. US2003018953A1  
; GENERAL INFORMATION:

; APPLICANT: Terry, Roemer D.  
; APPLICANT: Bo, Jiang  
; APPLICANT: Charles, Boone  
; APPLICANT: Howard, Bussey  
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery  
; FILE REFERENCE: 10182-005-999  
; CURRENT APPLICATION NUMBER: US/10/032,585  
; CURRENT FILING DATE: 2001-12-20  
; NUMBER OF SEQ ID NOS: 8000  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 2790  
; LENGTH: 65  
; TYPE: DNA  
; ORGANISM: Candida albicans  
US-10-032-585-2790

Query Match 100.0%; Score 10; DB 12; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 25 CTTCTCTTTT 34

## RESULT 98

US-10-015-637-3  
; Sequence 3, Application US/10015637  
; Publication No. US20030046727A1  
; GENERAL INFORMATION:

; APPLICANT: Wang, Qi  
; APPLICANT: Dubois, Patrice  
; APPLICANT: Liang, Jihong  
; APPLICANT: Oulmassov, Tim  
; TITLE OF INVENTION: Arcalin-5 Promoter and Uses Thereof  
; FILE REFERENCE: 13587.106  
; CURRENT APPLICATION NUMBER: US/10/015,637  
; CURRENT FILING DATE: 2001-12-17  
; PRIOR APPLICATION NUMBER: US 60/255879  
; PRIOR FILING DATE: 2000-12-18  
; NUMBER OF SEQ ID NOS: 14  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 3  
; LENGTH: 76  
; TYPE: DNA  
; ORGANISM: Glycine max  
US-10-015-637-3

Query Match 100.0%; Score 10; DB 14; Length 76;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||

Db 43 CTTCTCTTTT 52  
|||||

## RESULT 99

US-10-029-386-24504  
; Sequence 24504, Application US/10029386  
; Publication No. US20030194704A1  
; GENERAL INFORMATION:

; APPLICANT: Penn, Sharron G.  
; APPLICANT: Hanzel, David K.  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: AEOICA-X-2  
; CURRENT APPLICATION NUMBER: US/10/029,386  
; CURRENT FILING DATE: 2001-12-20  
; NUMBER OF SEQ ID NOS: 34288  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 24504  
; LENGTH: 81  
; TYPE: DNA  
; ORGANISM: Homo sapiens

; FEATURE:  
; OTHER INFORMATION: MAP TO CHR18.3  
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.1  
; OTHER INFORMATION: EST\_HUMAN HIT: AW905636.1, EVALUE 9.00e-02  
US-10-029-386-24504

Query Match 100.0%; Score 10; DB 12; Length 81;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||  
Db 29 CTTCTCTTTT 38

## RESULT 100

US-10-029-386-17114/c  
; Sequence 17114, Application US/10029386  
; Publication No. US20030194704A1  
; GENERAL INFORMATION:

; APPLICANT: Penn, Sharron G.  
; APPLICANT: Hanzel, David K.  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: AEOICA-X-2  
; CURRENT APPLICATION NUMBER: US/10/029,386  
; CURRENT FILING DATE: 2001-12-20  
; NUMBER OF SEQ ID NOS: 34288  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 17114  
; LENGTH: 84  
; TYPE: DNA  
; ORGANISM: Homo sapiens

; FEATURE:  
; OTHER INFORMATION: MAP TO AL137017.4  
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.7  
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.6  
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.6  
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.5  
; OTHER INFORMATION: EST\_HUMAN HIT: BE536158.1, EVALUE 9.50e-02  
; OTHER INFORMATION: NT HIT: X03248.1, EVALUE 1.00e+00  
US-10-029-386-17114

Query Match 100.0%; Score 10; DB 12; Length 84;  
Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10  
|||||



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Db 18 CTTCTTTT 9

Search completed: October 28, 2003, 18:49:54  
Job time : 1754 secs